

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
23 January 2003 (23.01.2003)

PCT

(10) International Publication Number
WO 03/006452 A1(51) International Patent Classification⁷: **C07D 311/04**,
311/32, 319/16, 327/06, 405/12, 411/12, C07C 217/14,
217/54, 217/90, A61K 31/135, 31/353, 31/357, 31/39,
31/4178, 31/4433, A61P 9/06, 9/10(21) International Application Number: **PCT/FI02/00621**(22) International Filing Date: **10 July 2002 (10.07.2002)**(25) Filing Language: **English**(26) Publication Language: **English**(30) Priority Data:
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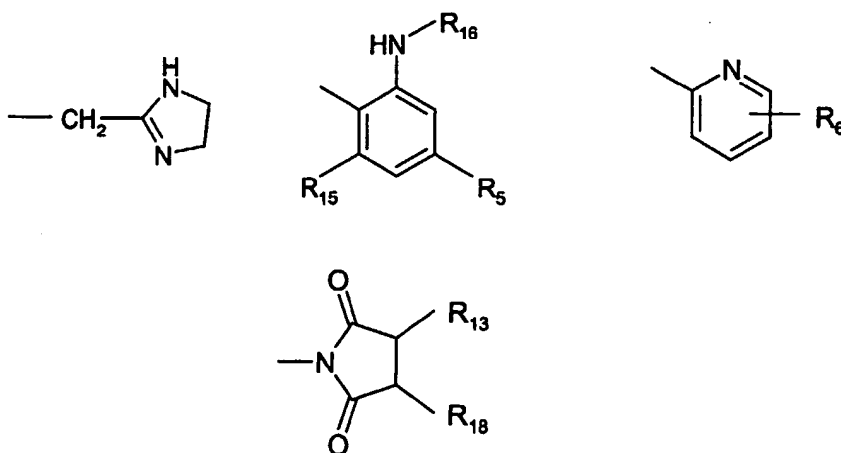
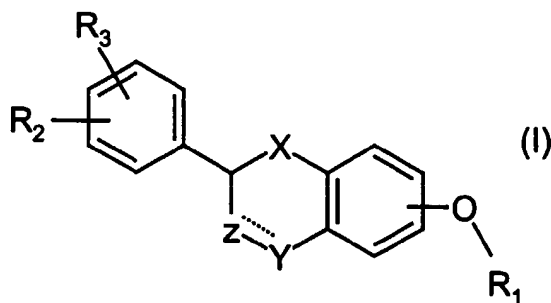
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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,

[Continued on next page]

(54) Title: **NEW COMPOUNDS, WHICH ARE POTENT INHIBITORS OF Na⁺/Ca²⁺ EXCHANGE MECHANISM AND ARE USEFUL IN THE TREATMENT OF ARRHYTHMIAS**

(57) Abstract: Therapeutically active compounds of formula (I): wherein X is -O-, -CH₂- or -C(O)-; Z is -CHR₉- or valence bond; Y is -CH₂-, -C(O)-, CH(OR₁₀)-, -CH(NR₁₁R₁₂)-, -O-, -S-, -S(O)- or -S(O₂)-, provided that in case Z is a valence bond, Y is not C(O); the dashed line represents an optional double bond in which case Z is -CR₉- and Y is -CH-, C(OR₁₀)- or -C(NR₁₁R₁₂)-; R₁ is -(CH₂)_nNR₄R₇ or one of the following groups: n is 1 - 4; R₂ and R₃ are independently H, lower alkyl, lower alkoxy, -NO₂, halogen, -CF₃, -OH, -NHR₈ or -COOH; R₄ and R₇ are independently H, lower alkyl or lower hydroxyalkyl; R₅ is H, lower alkoxy, -CF₃, -NH₂ or -CN; R₆ is -NO₂, -NR₁₄R₁₉, -CF₃ or R₈ and R₁₆ are independently H or acyl; R₉ is H or lower alkyl; R₁₀ is H, alkylsulfonyl or acyl; R₁₁ and R₁₂ are independently H, lower alkyl or acyl; R₁₃ and R₁₈ are independently H or -OR₂₀; R₁₄ and R₁₉ are independently H, acyl, alkylsulfonyl, C(S)NHR₁₇ or C(O)NHR₁₇; R₁₅ is H or NH₂; R₁₇ is H or lower alkyl; R₂₀ is H or acyl; and pharmaceutically acceptable salts and

esters thereof are disclosed. The compounds are potent inhibitors of Na⁺/Ca²⁺ exchange mechanism.

WO 03/006452 A1



MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,
VN, YU, ZA, ZM, ZW.

TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

(84) **Designated States (regional):** ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK,

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New compounds, which are potent inhibitors of $\text{Na}^+ / \text{Ca}^{2+}$ exchange mechanism and are useful in the treatment of arrhythmias

5 Technical field

The present invention relates to new therapeutically active compounds and pharmaceutically acceptable salts and esters thereof. The invention also relates to pharmaceutical compositions containing these compounds as active ingredients. The
10 compounds of the invention are potent inhibitors of $\text{Na}^+/\text{Ca}^{2+}$ exchange mechanism.

Background of the invention

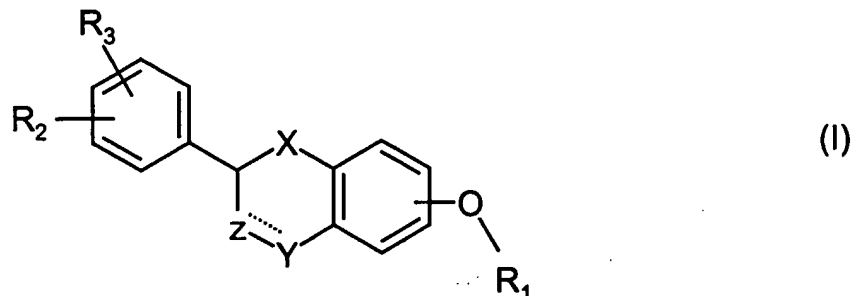
$\text{Na}^+/\text{Ca}^{2+}$ exchange mechanism is one of the ion transport mechanisms that regulate the concentration of sodium and calcium ions in the cells. Compounds which
15 selectively inhibit $\text{Na}^+/\text{Ca}^{2+}$ exchange mechanism and thereby prevent overload of Ca^{2+} in cells are regarded useful in preventing the cell injury mechanism of cardiac muscle and the like after ischemia and reperfusion. Such compounds are useful in the treatment of ischemic heart diseases, ischemic cerebral diseases, ischemic renal
20 diseases and in the protection of cells during thrombolytic therapy, angioplasty, bypass operation of coronary artery or organ transplantation and arrhythmias.

Compounds capable of inhibiting $\text{Na}^+/\text{Ca}^{2+}$ exchange system have been described earlier e.g. in patent publications WO 97/09306, EP 0978506, EP 1031556,
25 JP 11049752 and JP 11302235.

Summary of the invention

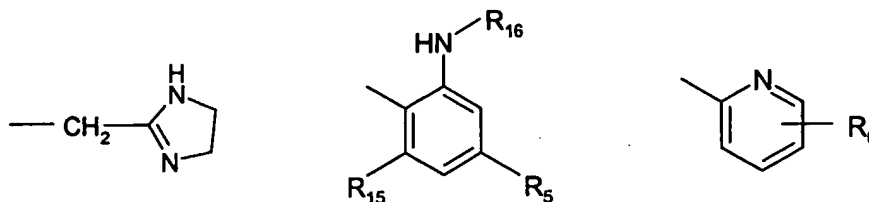
It has now been found that compounds of formula (I) are particularly potent
30 inhibitors of $\text{Na}^+/\text{Ca}^{2+}$ exchange mechanism and are particularly useful in the treatment of arrhythmias.

The compounds of the present invention have a structure represented by formula (I):



wherein

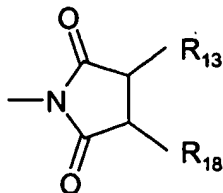
- 5 X is -O-, -CH₂- or -C(O)-;
 Z is -CHR₉- or valence bond;
 Y is -CH₂-, -C(O)-, CH(OR₁₀)-, -CH(NR₁₁R₁₂)-, -O-, -S-, -S(O)- or -S(O₂)-
 provided that in case Z is a valence bond, Y is not C(O);
 the dashed line represents an optional double bond in which case Z is -CR₉-
 10 and Y is -CH-, C(OR₁₀)- or -C(NR₁₁R₁₂)-;
 R₁ is -(CH₂)_nNR₄R₇ or one of the following groups:



n is 1 - 4,

R₂ and R₃ are independently H, lower alkyl, lower alkoxy, -NO₂, halogen, -CF₃, -OH, -NHR₈ or -COOH,

- 15 R₄ and R₇ are independently H, lower alkyl or lower hydroxyalkyl,
 R₅ is H, lower alkoxy, -CF₃, -NH₂ or -CN,
 R₆ is -NO₂, -NR₁₄R₁₉, -CF₃ or



R₈ and R₁₆ are independently H or acyl,

R₉ is H or lower alkyl,

- 20 R₁₀ is H, alkylsulfonyl or acyl;

R_{11} and R_{12} are independently H, lower alkyl or acyl,

R_{13} and R_{18} are independently H or $-OR_{20}$,

R_{14} and R_{19} are independently H, acyl, alkylsulfonyl, $C(S)NHR_{17}$ or

$C(O)NHR_{17}$,

5

R_{15} is H or NH_2 ,

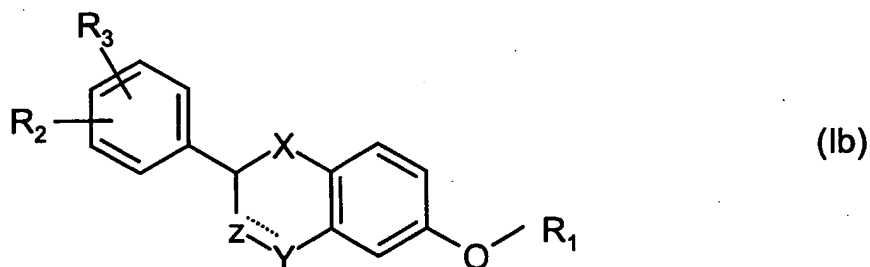
R_{17} is H or lower alkyl,

R_{20} is H or acyl,

and pharmaceutically acceptable salts and esters thereof.

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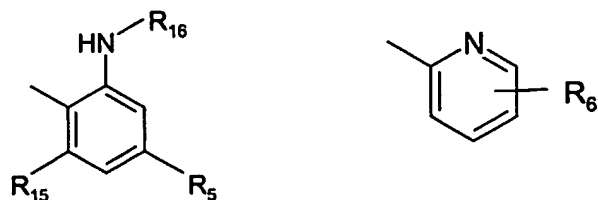
In one class of preferred compounds and pharmaceutically acceptable salts and esters thereof are compounds of formula



15 wherein R_1 , R_2 , R_3 , X, Y and Z are as defined above. In another class of preferred compounds and pharmaceutically acceptable salts and esters thereof are compounds of formula (I), wherein X is O, and Z and Y is $-CH_2-$. In another class of preferred compounds and pharmaceutically acceptable salts and esters thereof are compounds of formula (I), wherein X is O, and Z is $-CH_2-$ and Y is $CHOH$.

20

In one subclass of preferred compounds and pharmaceutically acceptable salts and esters thereof are compounds of formula (I) wherein R_1 is one of the following groups



In another subclass of preferred compounds are compounds, wherein R_6 is $-\text{NO}_2$ or $-\text{NR}_{14}\text{R}_{19}$. In a group of this subclass R_{14} and R_{19} are independently H, acyl or alkylsulfonyl, R_{15} and R_{16} is preferably H and R_5 is H or lower alkoxy.

5 In one subclass of preferred compounds are compounds, wherein R_2 and R_3 are independently H or halogen. Fluorine is the preferred halogen.

In one class of preferred compounds are compounds wherein $n = 2$. R_4 and R_7 are preferably methyl.

10

The present invention also provides a pharmaceutical composition comprising a compound of formula (I) together with a pharmaceutically acceptable carrier.

15 The present invention further provides a method for inhibiting $\text{Na}^+/\text{Ca}^{2+}$ exchange mechanism in a cell, comprising administering to a subject in need thereof a therapeutically effective amount of a compound of formula (I).

The present invention further provides a method for preventing overload of Ca^{2+} ions in cells, comprising administering to a subject in need thereof a
20 therapeutically effective amount of a compound of formula (I).

The present invention further provides a method for treating arrhythmias, comprising administering to a subject in need thereof a therapeutically effective amount of a compound of formula (I).

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Brief description of the drawings

FIG. 1 shows the antiarrhythmic effects of the compound of Example 27 on ouabain-induced aftercontractions in guinea-pig isolated papillary muscles.

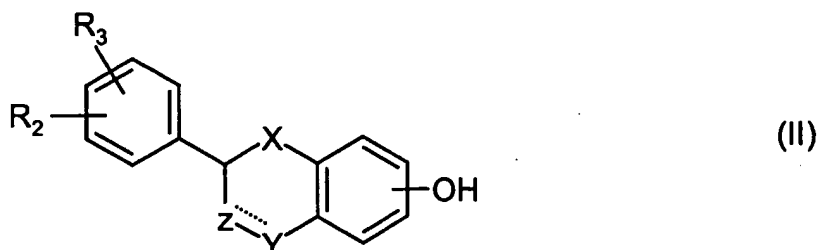
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FIG. 2 shows the antiarrhythmic effects of the compound of Example 67 on ouabain-induced aftercontractions in guinea-pig isolated papillary muscles.

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Detailed description of the invention

The compounds of the invention can be prepared from corresponding phenol derivatives (II), wherein R_2 , R_3 , X, Z and Y are the same as defined above.



5

The syntheses are shown in Scheme 1, wherein formula (II) is abbreviated as Ar-OH (II), and R_4 , R_5 , R_6 , R_7 , R_{15} and R_{16} are the same as defined above, and Hal means halogen.

10

Compounds of formula (II) can be reacted with 1-chloro-2-nitrobenzene derivatives to result in nitrophenoxy compounds (10), which in turn by hydrogenation gives phenylamine derivatives (11). 5-Nitropyridin-2-yloxy derivatives (12) are obtained by reactions with 2-chloro-5-nitropyridine.

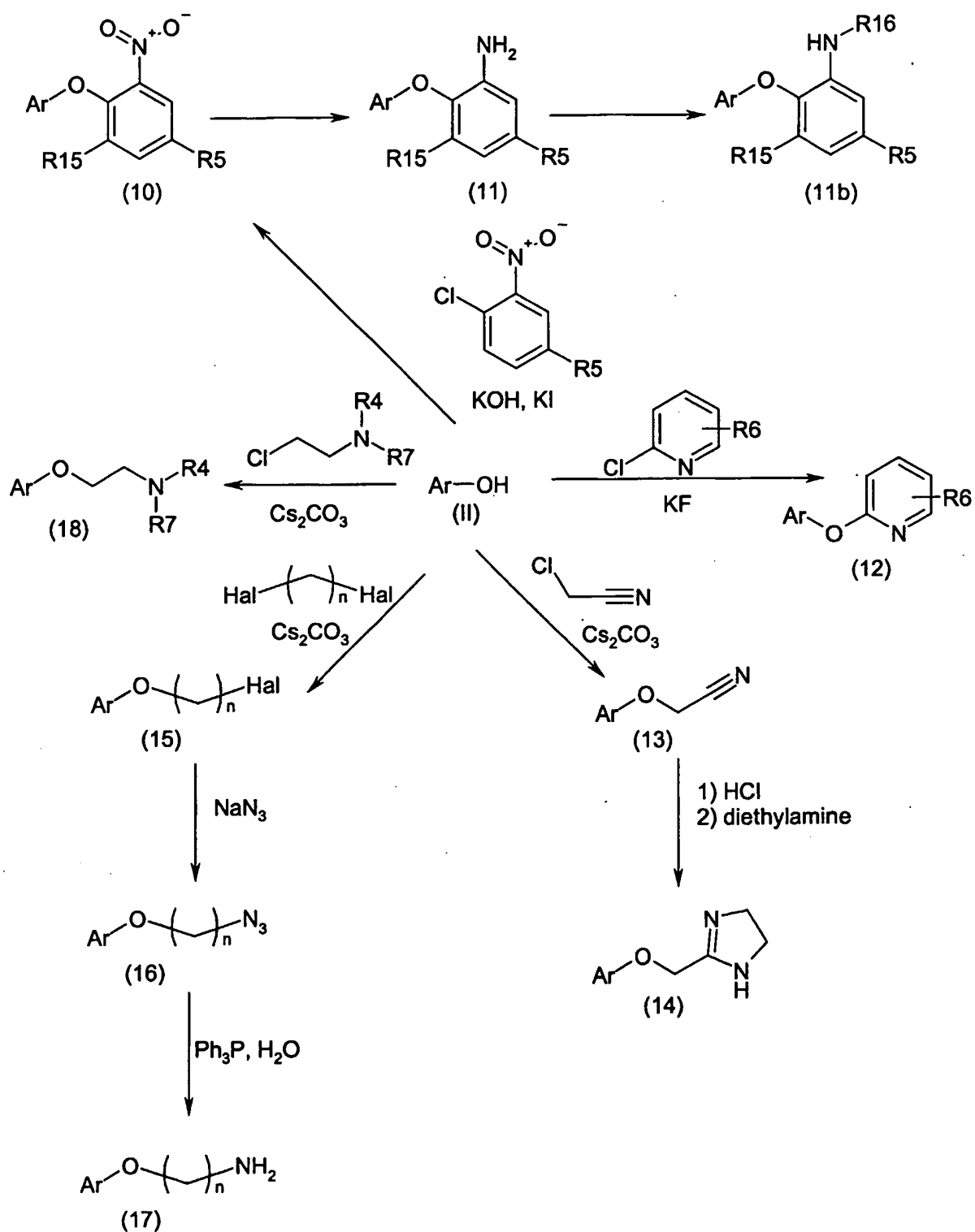
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2-Oxymethyl imidazoline derivatives (14) can be synthesised from phenol derivatives (II) via cyano methyl ether (13), which is converted to imidazoline (14) by a known method (e.g. *J. Med. Chem.* 1994, 37(12), 1874). Alkoxyazide derivatives (16) are obtained via corresponding haloalkoxy derivatives (15) by reaction with sodium azide. Azides (16) are converted to amines (17) by reaction with triphenylamine. 2-(Dimethylamino)ethoxy derivatives (18) are obtained by direct reactions of phenols of formula (II) with 2-(dimethylamino)ethyl chloride.

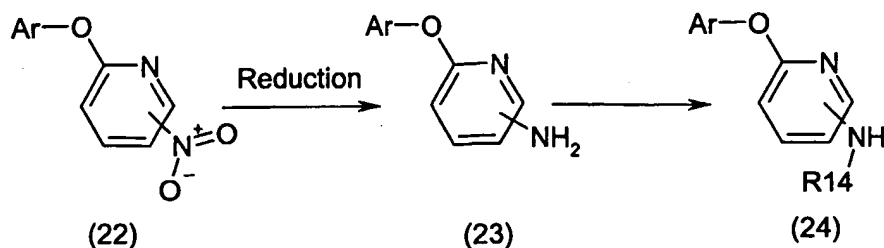
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The reduction of nitropyridines (22) followed by acylation, mesylation etc. produces compounds of formula (24), as shown by Scheme 2.

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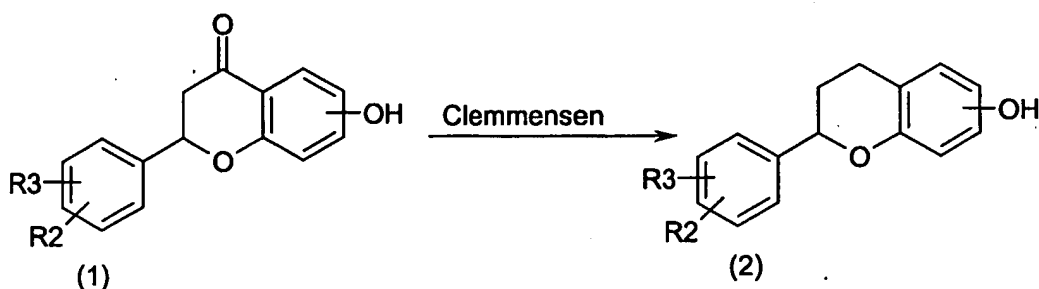
SCHEME 1.



SCHEME 2.

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As shown in the following Scheme 3, wherein R₂ and R₃ are the same as defined above, 6- and 7-hydroxyflavane derivatives (2) are obtained from corresponding flavanones (1) by Clemmensen reduction. 6- and 7-hydroxyflavanones (1) are commercially available or can be synthesised by methods described in the literature, e.g. *J. Org. Chem.*, 1960, 25, 1247-9 and *J. Org. Chem.*, 1958, 23, 1159-61 or as described later in Scheme 5.

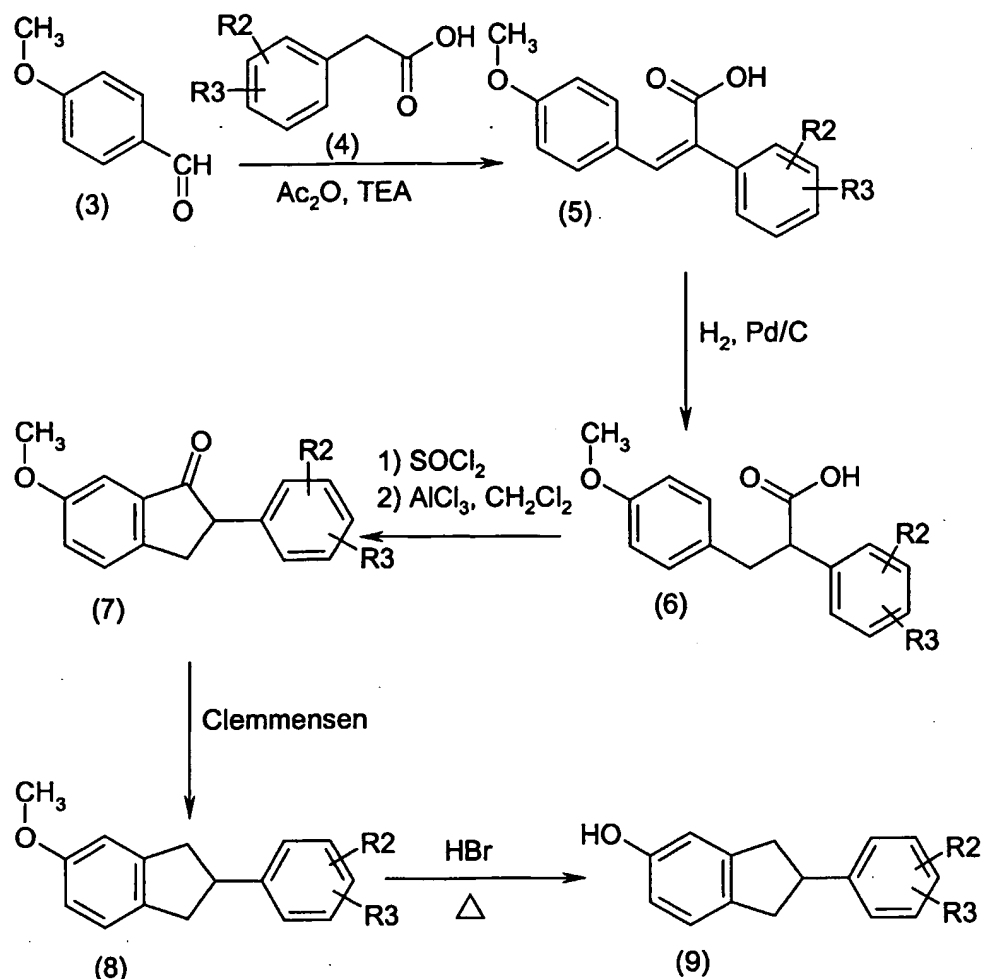


SCHEME 3.

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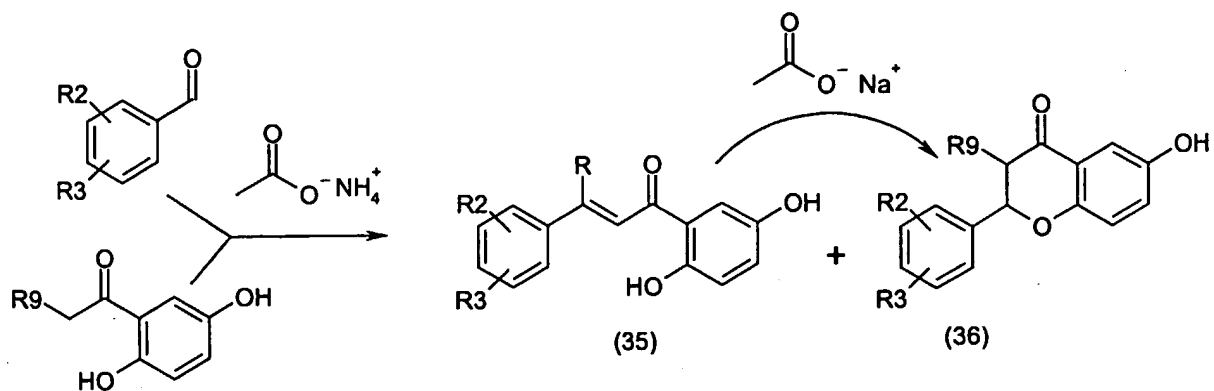
The following Scheme 4, wherein R₂ and R₃ are the same as defined above, describes the synthesis of 2-phenyl indan-5-ols (9). Condensation of *p*-anisaldehyde (3) with substituted phenyl acetic acid (4) gives mixture of *cis*- and *trans*-isomers of the corresponding acrylic acid (5). After hydrogenation and intramolecular Friedel-Crafts reaction carbonyl functionality of 1-indanones (7) can be reduced by Clemmensen reduction. Finally methoxy indane (8) is refluxed in concentrated hydrobromic acid to obtain 2-phenyl indan-5-ols (9).

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SCHEME 4

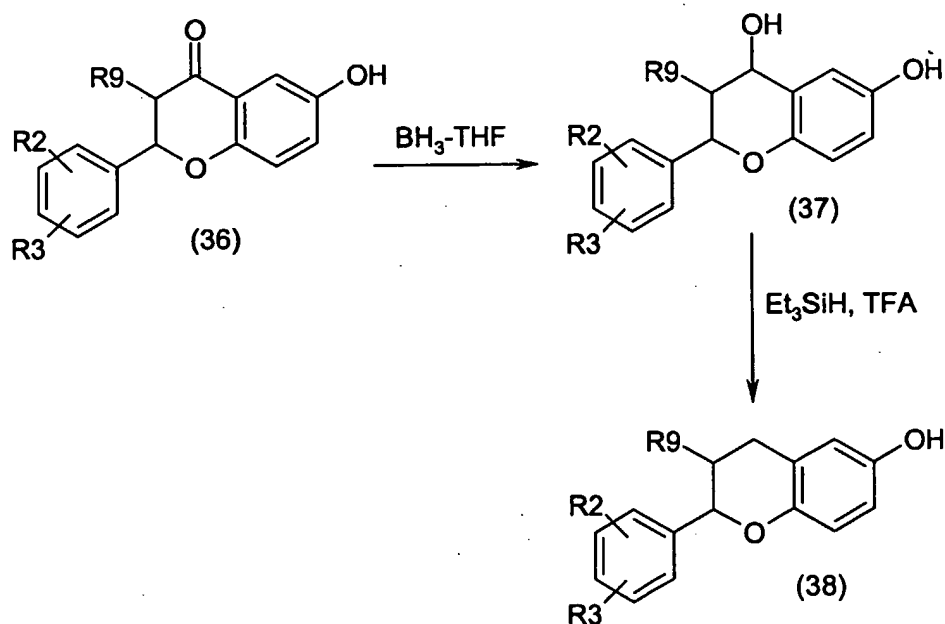
- 5 6-Hydroxyflavanone derivatives can be synthesised as shown in Scheme 5. 2',5'-Dihydroxyacetophenone or corresponding propiophenone is condensed with appropriate benzaldehyde resulting in a mixture of desired 6-hydroxyflavanone (36) and the corresponding chalcone (35). The chalcone can be cyclised to flavanone.



SCHEME 5

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2-Phenylchroman-4,6-diol derivatives (37) are obtained from corresponding 6-hydroxyflavanones (36) by reduction as shown in Scheme 6. These diol derivatives can be reduced further into 6-hydroxyflavanes (38).



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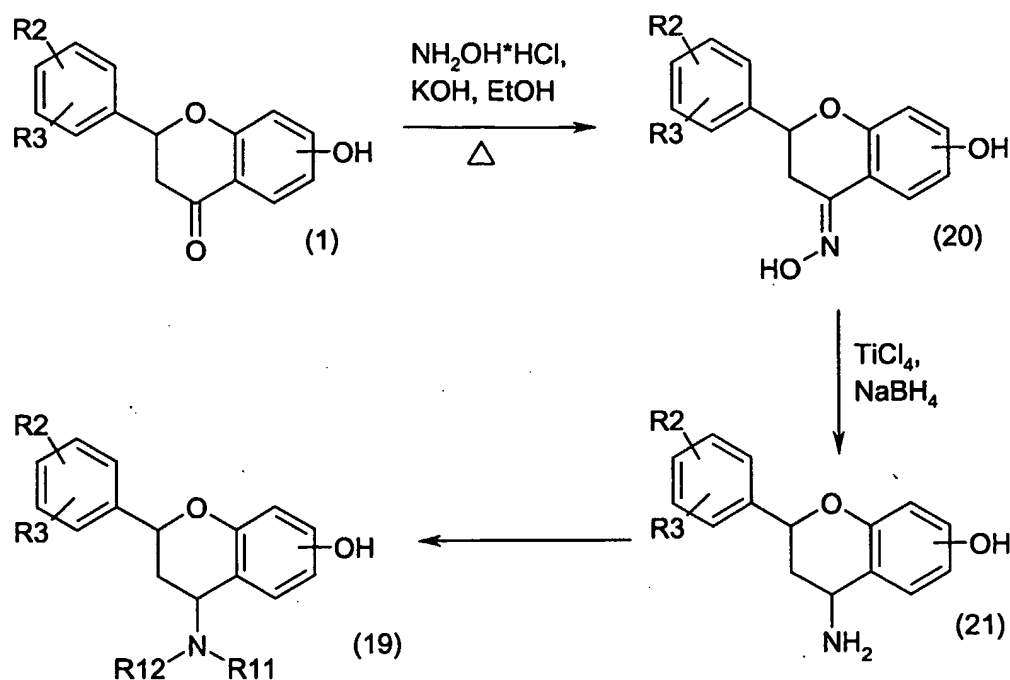
SCHEME 6

As shown in the following Scheme 7, the 4-amino-2-phenylchromanol derivatives (19) can be synthesised from corresponding flavanones (1) via hydroxy-2-

15

phenylchroman-4-one oximes (20). Reduction of oxime derivative gives 4-amino-2-phenylchromanols (21) which can be alkylated or acylated by common methods. 4-Hydroxy-2-phenylchromanol derivatives can be treated by similar methods.

5



SCHEME 7

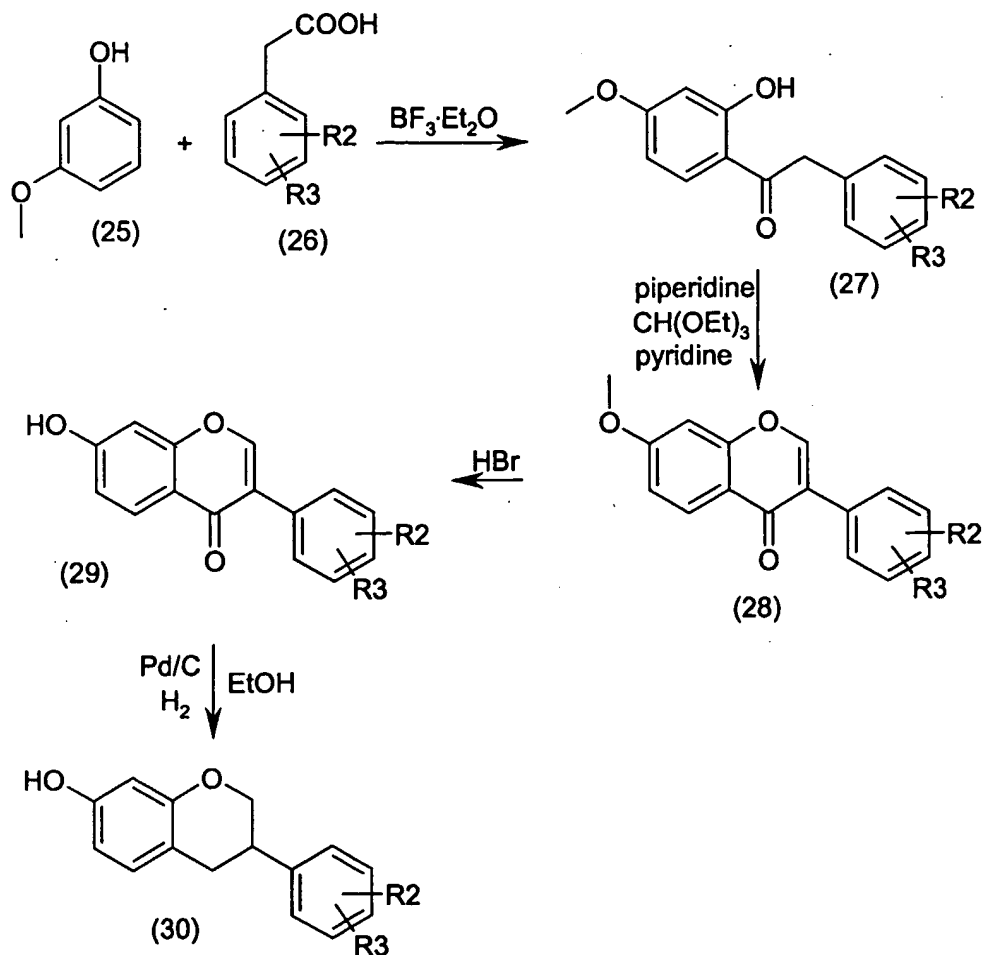
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The following Scheme 8, wherein R₂ and R₃ are the same as defined above, describes the synthesis of 7-hydroxyisoflavones (29) and 7-hydroxyisoflavans (30).

Acylation of 3-methoxyphenol with substituted phenyl acetic acids gives the corresponding 2-hydroxydeoxybenzoins (27) which can be cyclised with triethylorthoformate to yield isoflavones (28). Deprotection with hydrobromic acid and catalytic

15

hydrogenation gives 7-hydroxyisoflavans (30).

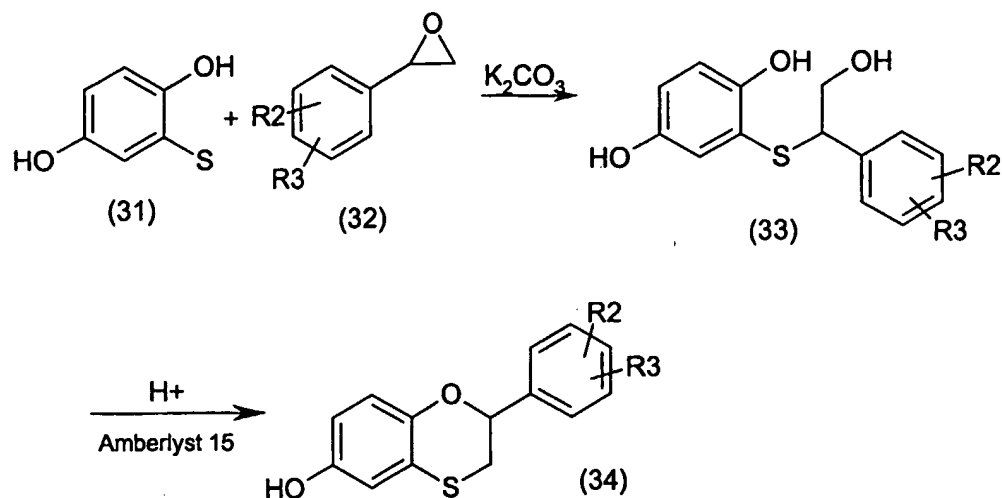


SCHEME 8.

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The following Scheme 9 describes the synthesis of 2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-ol (34). The reaction of 2-mercaptobenzene-1,4-diol with styrene epoxide in the presence of base gives sulfide (33). The ring closure with an acid ion exchanger affords 2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-ol (34).

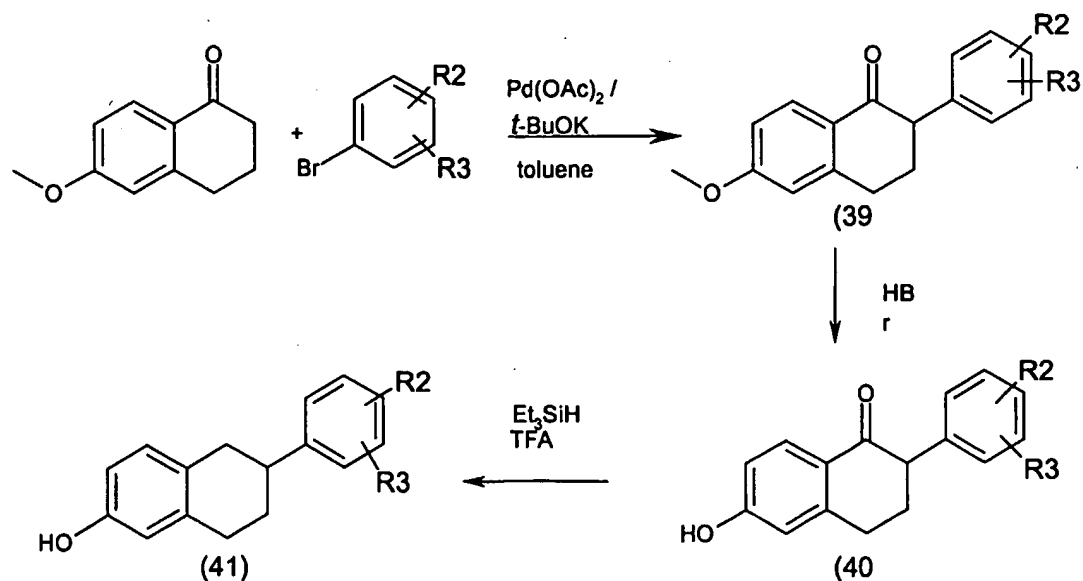
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SCHEME 9.

5 The following Scheme 10 describes the synthesis of 6-phenyl-5,6,7,8-tetrahydro-naphthalen-2-ol (41) and 6-hydroxy-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one (40). Pd-catalyzed α -arylation of 6-methoxy-1-tetralone gives 6-methoxy-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one (39) which after demethylation leads to the phenolic compound (40).

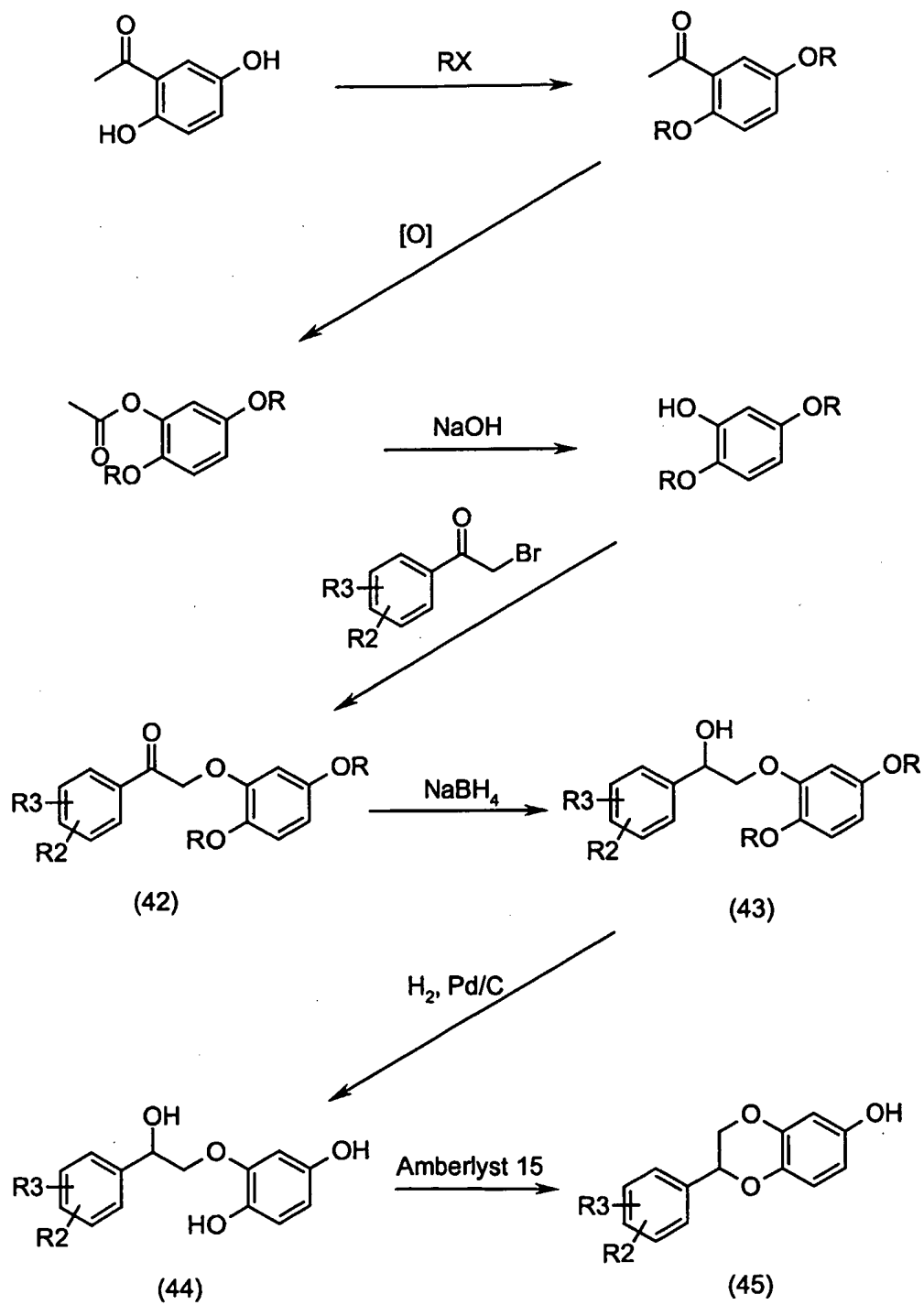
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SCHEME 10

The following Scheme (11), wherein R_2 and R_3 are the same as defined above, describes the synthesis of 2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-ols (45). After protecting hydroxyl groups of 2,5-dihydroxyacetophenone this ketone rearranges with peracids and gives a phenol after hydrolysis. The phenol is condensed with a
5 haloketone and after reduction and removal of protection groups the hydroxyphenol (44) is cyclised to a 2,3-dihydro-2-phenyl-benzo[1,4]-dioxin-6-ol (45).

14



SCHEME 11

Salts and esters of the compounds, when applicable, may be prepared by known methods. Physiologically acceptable salts are useful as active medicaments. Examples are the salts with inorganic acids such as hydrochloric acid, hydrobromic acid or nitric acid, and salts with organic acids such as methanesulfonic acid, citric acid or tartaric acid. Physiologically acceptable esters are also useful as active medicaments. Examples are the esters with aliphatic or aromatic acids such as acetic acid or with aliphatic or aromatic alcohols.

The term "alkyl" as employed herein by itself or as part of another group includes both straight, branched and cyclized chain radicals of up to 18 carbon atoms, preferably 1 to 7 carbon atoms, most preferably 1 to 4 carbon atoms. The term "lower alkyl" as employed herein by itself or as part of another group includes straight, branched and cyclized chain radicals of 1 to 7, preferably 1 to 4, most preferably 1 or 2 carbon atoms. Specific examples for the alkyl and lower alkyl residues, respectively, are methyl, ethyl, propyl, isopropyl, butyl, tert. butyl, pentyl, cyclopentyl, hexyl, cyclohexyl, octyl, decyl and dodecyl including the various branched chain isomers thereof.

The term "alkoxy" as employed herein by itself or as part of another group includes an alkyl group as defined above linked to an oxygen atom.

The term "acyl" as employed herein by itself or as part of another group refers to an alkylcarbonyl or alkenylcarbonyl group, the alkyl and alkenyl groups being defined above.

Compounds of the invention may be administered to a patient in therapeutically effective amounts which range usually from about 0.05 to 200 mg, preferably 0.1 to 100 mg, more preferably 0.5 to 50, mg per day depending on the age, weight, condition of the patient, administration route and the $\text{Na}^+/\text{Ca}^{2+}$ exchange inhibitor used. The compounds of the invention can be formulated into dosage forms using the principles known in the art. It can be given to a patient as such or in combination with suitable pharmaceutical excipients in the form of tablets, granules, capsules, suppositories, emulsions, suspensions or solutions. Choosing suitable ingredients for the composition is a routine for those of ordinary skill in the art. It is evident that suitable carriers, solvents, gel forming ingredients, dispersion forming ingredients, antioxidants, colours, sweeteners, wetting compounds and other ingredients normally

used in this field of technology may be also used. The compositions containing the active compound can be given enterally or parenterally, the oral route being the preferred way. The contents of the active compound in the composition is from about 0.5 to 100 %, preferably from about 0.5 to about 20 %, per weight of the total composition.

EXPERIMENTS

The effects of the compounds of the invention were tested on ouabain-induced arrhythmias in guinea-pig papillary muscles.

Methods

Guinea-pig papillary muscles were mounted into horizontal muscle cuvette. A hook connected to force transducer was attached to another end of the muscle. Muscle preparations were electrically paced at 1 Hz with field stimulation via platinum electrodes. Modified Tyrode solution was used for superfusion of muscle preparations. The composition of the Tyrode solution was the following (mM): NaCl 135, MgCl₂ x 6H₂O 1, KCl 5, CaCl₂ x 2H₂O 2, NaHCO₃ 15, Na₂HPO₄ x 2H₂O 1, and glucose 10. The Tyrode solution was gassed with carbogen (95% O₂, 5% CO₂) to set pH at 7.4. Experiments were carried out at 37 °C. Acquisition and analysis of twitch tensions with Action Potential and Force Measurement System (ACFO v1.0, Fision Ltd, Finland).

Inhibition of ouabain-induced arrhythmias

Ouabain by blocking of sodium-potassium ATPase increase intracellular sodium which is changed for calcium via NCX. Increased intracellular calcium is leading to overload of sarcoplasmic reticulum (SR) and spontaneous calcium release from SR inducing delayed afterpolarizations (DADs). Equivalence for DADs in force signal is aftercontractions (ACs) which are seen as spontaneous twitches after the pacing controlled twitch.

The compounds of the invention delayed appearance and decreased amplitude of aftercontractions. As shown in Figure 1, the title compound of Example 27, at 30 µM concentration, delayed appearance [38 ± 7.5 min vs vehicle: 25 ± 8.9 min (mean

± SD), $p=0.013$, $n=5$] and decreased maximum amplitude of aftercontractions #1 (74 ± 16 mg vs vehicle: 143 ± 54 mg, $p=0.008$, One-way ANOVA followed by LSD; $n=5$). As shown in Figure 2, the title compound of Example 67, at $10 \mu\text{M}$, decreased maximum amplitude of aftercontractions #1 (88 ± 20 mg vs vehicle: 143 ± 54 mg, $p=0.027$, $n=5$).

EXAMPLES:

Example 1:

5-Nitro-2-(2-phenylchroman-6-yloxy)pyridine

a) 2-phenylchroman-6-ol

Zinc (5,4 g, 83,2 mmol), mercury (II) chloride (340 mg), concentrated hydrogen chloride (0,2 ml) and water were mixed at room temperature for 15 minutes and the mixture was decanted. 6-Hydroxyflavanone (1,0 g) was added as a suspension in a mixture of acetic acid (25 ml), concentrated hydrogen chloride (5,2 ml) and water (2 ml). The reaction mixture was refluxed for $1\frac{1}{2}$ hours. After cooling into room temperature, the reaction mixture was filtered and the filtrate was extracted with ethyl acetate. The combined organic layers were washed with saturated NaHCO_3 -solution, then with water and dried with Na_2SO_4 . The 2-phenylchroman-6-ol was purified by column chromatography using heptane-ethyl acetate (2:1) as an eluant. ^1H NMR (400 MHz, d_6 -DMSO) δ : 8.78 (s, 1H), 7.43-7.31 (m, 5H), 6.63 (d, 1H, J 8.6 Hz) 6.51 (dd, 1H, J 8.6, 2.9 Hz), 6.48 (d, 1H, J 2.9 Hz), 4.98 (dd, 1H, J, 9.9, 2.2 Hz), 2.89 (ddd, 1H, J -16.7, 11.3, 6.1 Hz), 2.63 (ddd, 1H, J -16.7, 5.5, 3.3 Hz) 2.10 (m, 1H), 1.94 (m, 1H).

b) 5-Nitro-2-(2-phenylchroman-6-yloxy)pyridine

Potassium fluoride (225 mg) was added into a solution of 2-phenylchroman-6-ol (300 mg) in dry DMF (3 ml). After stirring the resulting mixture at 120°C for 30 minutes 2-chloro-5-nitropyridine (195 mg) was added. The reaction mixture was stirred for a further $6\frac{1}{2}$ hours at 120°C . After cooling into room temperature 1 M HCl-solution was added and the mixture was extracted with ethyl acetate. The combined organic layers were washed with water then with saturated NaCl-solution and dried with Na_2SO_4 . 5-Nitro-2-(2-phenylchroman-6-yloxy)-pyridine was

recrystallised from acetone- 2-propanol (1:5). ¹H NMR (400 MHz, d₆-DMSO) δ: 9.00 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.2, 2.9 Hz), 7.47-7.32 (m, 5H), 7.20 (d, 1H, J 9.2 Hz), 7.00-6.89 (m, 3H), 5.15 (dd, 1H, J 10.1, 2.2 Hz), 2.99 (ddd, 1H, J -16.8, 11.3, 6.2 Hz), 2.75 (ddd, 1H, J -16.8, 5.4, 3.3 Hz) 2.18 (m, 1H), 2.02 (m, 1H).

5

Example 2:Dimethyl[2-(2-phenylchroman-6-yloxy)ethyl]amine

Cesium carbonate (230 mg) and an excess of 2-(dimethylamino)ethyl chloride
10 in ethyl acetate were added into a solution of 2-phenylchroman-6-ol (150 mg) in acetonitrile (5 ml). The reaction mixture was refluxed for 30 minutes. After removing the solvents, the residue was taken up in water and extracted with ethyl acetate. The combined organic layers were washed with water and then with saturated NaCl-solution and dried with Na₂SO₄. The dimethyl[2-(2-phenylchroman-6-yloxy)-
15 ethyl]amine was crystallised from heptane. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.43-7.30 (m, 5H) 6.75-6.67 (m, 3H), 5.03 (dd, 1H, J 10.0, 2.2 Hz), 3.95 (t, 2H, J 5.9 Hz), 2.94 (ddd, 1H, J -16.7, 10.9, 5.8 Hz), 2.69 (ddd, 1H, J -16.7, 5.2, 3.3 Hz), 2.58 (t, 2H, J 5.9 Hz) 2.13 (m, 1H), 1.96 (m, 1H).

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Example 3:5-Methoxy-2-(2-phenyl-chroman-6-yloxy)phenylamine hydrochloride

a) 6-(4-Methoxy-2-nitrophenoxy)-2-phenylchroman

The 2-phenylchroman-6-ol (500 mg) and 1-chloro-4-methoxy-2-nitrobenzene
25 (390 mg) were dissolved in DMSO (10 ml). Potassium hydroxide (230 mg) and potassium iodide (520 mg) were added and the resulting mixture was stirred at 90°C for 1 hour. After cooling it was poured in to 1 M HCl-solution (20 ml) and extracted with dichloromethane. The combined organic layers were washed with water until
30 neutral and then with saturated NaCl-solution and dried with Na₂SO₄. After evaporating the solvents the 6-(4-Methoxy-2-nitrophenoxy)-2-phenylchroman was obtained by trituration with methanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.58 (d, 1H, J 3.1 Hz), 7.45-7.33 (m, 6H), 7.28 (dd, 1H, J 9.2 Hz, 3.1 Hz) 7.10 (d, 1H, J 9.2 Hz), 6.86-6.78 (m, 3H), 5.10 (dd, 1H, J 10.0, 1.9 Hz), 3.83 (s, 3H) 2.92 (ddd, 1H, J -16.9, 11.2,
35 5.9 Hz), 2.75 (ddd, 1H, J -16.9, 7.9, 4.2 Hz) 2.15 (m, 1H), 1.97 (m, 1H).

b) 5-Methoxy-2-(2-phenyl-chroman-6-yloxy)phenylamine hydrochloride

6-(4-Methoxy-2-nitrophenoxy)-2-phenylchroman (360 mg) was dissolved in ethyl acetate and 10 % palladium on carbon (90 mg) was added. The reaction mixture was hydrogenated for 2 hours at normal pressure and room temperature. It was then filtered through Celite and washed with ethyl acetate. The 5-methoxy-2-(2-phenyl-chroman-6-yloxy)phenylamine was isolated as its hydrochloride salt. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.44-7.33 (m, 5H) 6.83-6.72 (m, 4H), 6.68 (d, 1H, J 2.9 Hz), 6.48 (dd, 1H, J 8.9, 2.8 Hz) 5.07 (dd, 1H, J 10.0, 2.2 Hz), 3.70 (s, 3H) 2.93 (ddd, 1H, J -17.0, 11.1, 6.1 Hz), 2.68 (ddd, 1H, J -17.0, 8.3, 4.5 Hz) 2.15 (m, 1H), 1.97 (m, 1H).

Example 4:

2-(2-Phenylchroman-6-yloxy)ethylamine methane sulfonate

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a) 6-(2-azidoethoxy)-2-phenylchroman

2-Phenylchroman-6-ol (340 mg), 1-bromo-2-chloroethane (1.25 ml) and cesium carbonate (977 mg) were dissolved in acetonitrile (4 ml). The reaction mixture was refluxed for 4 hours. After cooling into room temperature it was poured in 1 M HCl-solution and extracted with dichloromethane. The combined dichloromethane extracts were washed with water and dried with Na₂SO₄. The mixture was passed through silica gel column using ethyl acetate- hexane (1:7) as an eluant resulting 190 mg of haloethane derivative. It was dissolved in DMF (5 ml) and sodium azide (214 mg) was added. The reaction mixture was refluxed for 2 hours. The mixture was filtered. Ethyl acetate was added to the filtrate and it was then washed once with 1 M HCl-solution and then several times with water and dried with Na₂SO₄. The solvents were evaporated under reduced pressure to give 6-(2-azidoethoxy)-2-phenylchroman. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.44-7.32 (m, 5H) 6.78-6.71 (m, 3H), 5.04 (dd, 1H, J 10.1, 2.3 Hz), 4.10 (t, 2H, J 4.8 Hz), 3.60 (t, 2H, J 4.8 Hz), 2.95 (ddd, 1H, J -16.8, 11.1, 6.0 Hz), 2.70 (ddd, 1H, J -16.8, 5.3, 3.3 Hz), 2.14 (m, 1H), 1.97 (m, 1H).

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b) 2-(2-Phenylchroman-6-yloxy)ethylamine methane sulfonate

Triphenylphosphine (165 mg) and 40 µl of water were added into a solution of 6-(2-azidoethoxy)-2-phenylchroman (155 mg) in tetrahydrofuran. The resulting mixture was stirred for 2 hours at room temperature. 2-(2-Phenylchroman-6-yloxy)-ethylamine was isolated as its methane sulfonate salt. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.91 (bs, 3H), 7.44-7.32 (m, 5H), 6.81-6.75 (m, 3H), 5.05 (dd, 1H, J 9.9, 2.3 Hz), 4.08 (t, 2H, J 5.1 Hz), 3.19 (m, 2H), 2.95 (ddd, 1H, J -16.8, 11.0, 5.9. Hz), 2.71 (ddd, 1H, J -16.8, 5.2, 3.4 Hz), 2.30 (s, 3H) 2.15 (m, 1H), 1.97 (m, 1H).

Example 5:

2-(2-Phenylchroman-6-yloxymethyl)-4,5-dihydro-1H-imidazole hydrochloride

a) (2-phenylchroman-6-yloxy)acetonitrile

Cesium carbonate (310 mg) and chloroacetonitrile (62 µl) were added into a solution of 2-phenylchroman-6-ol (200 mg) in acetonitrile (3 ml). The resulting mixture was refluxed for 6 hours. The reaction mixture was allowed to cool to room temperature and 1 M HCl-solution was added and it was extracted with ethyl acetate. The combined organic layers were washed with water and saturated NaCl-solution and dried with Na₂SO₄. The solvents were evaporated under reduced pressure to give (2-phenylchroman-6-yloxy)acetonitrile. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.44-7.30 (m, 5H), 6.86-6.81 (m, 3H), 5.08 (dd, 1H, J 9.8, 2.2 Hz), 5.07 (s, 2H), 2.97 (ddd, 1H, J -16.9, 10.9, 6.0. Hz), 2.71 (ddd, 1H, J -16.9, 5.0, 3.4 Hz), 2.15 (m, 1H), 1.97 (m, 1H).

b) 2-(2-Phenylchroman-6-yloxymethyl)-4,5-dihydro-1H-imidazole hydrochloride

Dry HCl was passed through a solution of (2-phenylchroman-6-yloxy)acetonitrile (270 mg) in diethyl ether and 90 µl of absolute ethanol while cooling with ice bath. Reaction mixture was evaporated to the dryness after formation of the intermediate imidate. Precipitate was dissolved in absolute ethanol and 252 µl ethylene diamine was added to the cooled solution. Reaction mixture was allowed to warm to room temperature, evaporated to dryness, dissolved in dichloromethane and washed with water. Combined organic layers were dried and treated with charcoal. The 2-(2-Phenylchroman-6-yloxymethyl)-4,5-dihydro-1H-imidazole was isolated as its HCl-salt. ¹H-NMR (d₄-MeOH): 7.5-7.2 (m, 5H), 6.85-6.75 (m, 3H), 5.01 (d, 1H,

J 8.8 Hz), 4.97 (s, 2H), 4.00 (s, 4H), 3.02-2.90 (m, 1H), 2.80-2.70 (m, 1H), 2.21-2.12 (m, 1H), 2.05-1.90 (m, 1H). (M)⁺ = 308 (100%)

Example 6:**5 6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-one**

6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-one was prepared as described for 5-Nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 200 mg of 6-hydroxyflavanone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.03 (bs, 1H), 8.64 (d, 1H, J 9.0 Hz), 7.59-7.41 (m, 7H), 7.31 (d, 1H, J 9.0 Hz), 7.23 (d, 1H, 8.8 Hz), 5.75 (dd, 1H, J 12.3, 2.9 Hz), 3.30 (dd, 1H, -16.3, 12.3 Hz), 2.87 (dd, 1H, -16.3, 2.9 Hz).

Example 7:**15 7-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-one**

7-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-one was prepared as described for 5-Nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 150 mg of 7-hydroxyflavanone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.07 (d, 1H, J 2.8 Hz), 8.67 (dd, 1H, J 9.0, 2.8 Hz), 7.89 (d, 1H, 8.6 Hz), 7.60-7.35 (m, 6H), 7.04 (d, 1H, 2.1 Hz), 6.97 (dd, 1H, 8.6, 2.1 Hz), 5.75 (dd, 1H, J 13.0, 2.7 Hz), 3.32 (dd, 1H, 16.9, 13.0 Hz), 2.85 (d, -16.9, 2.7 Hz).

Example 8:**25 6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ol****a) 2-Phenylchroman-4,6-diol**

Into a suspension of 6-hydroxyflavanone (1.0 g) in dry THF (11.5 ml) was added dropwise a solution of borane-THF complex (12.5 ml, 1.0 M in THF) under nitrogen. The reaction mixture was refluxed for 1 hour. After cooling to the room temperature it was poured into an ice-2 M HCl-solution. 2-Phenylchroman-4,6-diol was filtered. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.83 (s, 1H), 7.45-7.38 (m, 4H), 7.35 (m, 1H), 6.89 (d, 1H, J 2.8 Hz), 6.59 (d, 1H, J 8.7 Hz), 6.54 (dd, 1H, J 8.7, 2.8 Hz), 5.41 (d, 1H, J 7.0 Hz), 5.11 (dd, 1H, J 11.7, 1.2 Hz), 4.87 (m, 1H), 2.26 (m, 35 1H), 1.90 (m, 1H).

b) 6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ol

6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 1.5 g of 2-phenylchroman-4,6-diol. The product was passed through a silica gel column using toluene - ethyl acetate (4:1) as an eluant and then crystallised from 2-propanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.7 Hz), 8.61 (dd, 1H, J 9.1, 2.7 Hz), 7.50-7.36 (m, 5H), 7.25 (d, 1H, J 2.7 Hz), 7.22 (d, 1H, J 9.1 Hz), 7.00 (dd, 1H, J 8.7, 2.7 Hz), 6.88 (d, 1H, J 8.7 Hz), 5.65 (d, 1H, J 6.3 Hz), 5.30 (dd, 1H, J 11.9, 1.3 Hz), 4.99 (m, 1H), 2.33 (m, 1H), 1.98 (m, 1H).

Example 9:2-[2-(3-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(3-Fluorophenyl)-6-hydroxychroman-4-one

2',5'-Dihydroxyacetophenone (1.50 g) was dissolved in warm glacial acetic acid (26 ml). 3-Fluorobenzaldehyde (1.35 g) and ammonium acetate (0.98 g) were added. The reaction mixture was refluxed for 2 hours. It was allowed to cool to room temperature and poured in ice. The precipitate formed was filtered resulting in 2.2 g of a mixture of 2-(3-fluorophenyl)-6-hydroxychroman-4-one and 1-(2,5-dihydroxyphenyl)-3-(3-fluorophenyl)propenone. The obtained mixture was dissolved in ethanol (90 ml) and sodium acetate (1.75 g) was added. The reaction mixture was refluxed for 5 hours. It was then allowed to cool to room temperature and diluted with water and filtered. The 2-(3-fluorophenyl)-6-hydroxychroman-4-one was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.45 (s, 1H), 7.47 (m, 1H), 7.40-7.37 (m, 2H), 7.22 (m, 1H), 7.12 (d, 1H, J 3.0 Hz), 7.05 (dd, 1H, J 8.8, 3.0 Hz), 6.98 (d, 1H, J 8.8 Hz), 5.59 (dd, 1H, J 13.0, 2.9 Hz), 3.21 (dd, 1H, J -16.9, 13.0 Hz), 2.82 (dd, 1H, J -16.9, 2.9 Hz).

b) 2-(3-Fluorophenyl)chroman-4,6-diol

2-(3-Fluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 220 mg of 2-(3-fluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 7.45 (m, 1H), 7.30-7.25 (m, 2H), 7.15 (m, 1H), 6.88 (d, 1H, J 2.8 Hz), 6.62 (d, 1H, J 8.7 Hz),

6.55 (dd, 1H, J 8.7, 2.8 Hz), 5.44 (d, 1H, J 7.0 Hz), 5.15 (d, 1H, J 10.7 Hz), 4.86 (m, 1H), 2.29 (m, 1H), 1.86 (m, 1H).

c) 2-(3-Fluorophenyl)chroman-6-ol

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Triethylsilane (960 μ l) was added slowly into a solution of 2-(3-fluorophenyl)chroman-4,6-diol (195 mg) in dichloromethane (4 ml). Trifluoroacetic acid (1.9 ml) was then added dropwise into a reaction mixture and it was stirred at room temperature for 5 hours. The reaction mixture was poured on ice-water and extracted with dichloromethane. The residue was evaporated under reduced pressure with toluene to obtain 2-(3-fluorophenyl)chroman-6-ol. ^1H NMR (400 MHz, d_6 -DMSO) δ : 8.78 (s, 1H), 7.43 (m, 1H), 7.28-7.25 (m, 2H), 7.14 (m, 1H), 6.66 (d, 1H, J 8.5 Hz) 6.52 (dd, 1H, J 8.5, 2.7 Hz), 6.49 (d, 1H, J 2.7 Hz), 5.03 (dd, 1H, J 9.9, 2.1 Hz), 2.86 (m, 1H), 2.63 (m, 1H) 2.13 (m, 1H), 1.93 (m, 1H).

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d) 2-[2-(3-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(3-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 210 mg of 2-(3-fluorophenyl)chroman-6-ol. The product was recrystallised from 2-propanol. ^1H NMR (400 MHz, CDCl_3) δ : 9.07 (d, 1H, J 2.8 Hz), 8.46 (dd, 1H, J 9.0, 2.8 Hz), 7.36 (m, 1H), 7.21-7.15 (m, 2H), 7.03 (m, 1H), 7.01 (d, 1H, J 9.0 Hz), 6.98 (d, 1H, J 8.6 Hz) 6.92 (dd, 1H, J 8.6, 2.7 Hz), 6.90 (d, 1H, J 2.7 Hz), 5.09 (dd, 1H, J 10.3, 2.4 Hz), 3.01 (ddd, 1H, J -16.9, 11.4, 6.0 Hz), 2.82 (ddd, 1H, J -16.9, 5.1, 3.2 Hz) 2.24 (m, 1H), 2.09 (m, 1H).

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Example 10:

5-Nitro-2-(2-phenylchroman-7-yloxy)pyridine

30 a) 2-Phenylchroman-7-ol

2-Phenyl-chroman-7-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(a) starting from 1.0 g of 7-hydroxy-flavanone. The product was purified by column chromatography using heptane-ethyl acetate (2:1) as an eluant. ^1H NMR (400 MHz, CD_3OD) δ : 7.41-7.28 (m, 5H), 6.86

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(d, 1H, J 8.2 Hz) 6.32 (dd, 1H, J 8.2, 2.4 Hz), 6.29 (d, 1H, J 2.4 Hz), 5.00 (dd, 1H, J 9.9, 2.4 Hz), 2.84 (m, 1H), 2.64 (m, 1H) 2.15 (m, 1H), 1.99 (m, 1H).

b) 5-Nitro-2-(2-phenylchroman-7-yloxy)pyridine

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5-Nitro-2-(2-phenylchroman-7-yloxy)pyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 115 mg of 2-phenyl-chroman-7-ol. The product was purified on preparative TLC-plate covered with silica gel using toluene - ethyl acetate (15:1) as an eluant. ¹H NMR (400
10 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.8 Hz), 8.60 (dd, 1H, J 9.1, 2.8 Hz), 7.46-7.32 (m, 5H), 7.22 (d, 1H, J 9.1 Hz), 7.20 (d, 1H, J 8.9 Hz) 6.72 (dd, 1H, J 8.9, 2.3 Hz), 6.72 (d, 1H, J 2.3 Hz), 5.16 (dd, 1H, J 10.1, 2.1 Hz), 2.97 (ddd, 1H, J -16.7, 11.3, 5.9 Hz), 2.77 (ddd, 1H, J -16.7, 8.1, 4.5 Hz) 2.20 (m, 1H), 2.02 (m, 1H).

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Example 11:

2-[2-(2,4-Dichlorophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(2,4-Dichlorophenyl)-6-hydroxychroman-4-one

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2-(2,4-Dichlorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 1.0 g of 2',5'-dihydroxyacetophenone and 1.4 g of 2,4-dichlorobenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.49 (s, 1H), 7.78 (d, 1H, J 8.5 Hz), 7.71(d, 1H, J 2.0 Hz)), 7.57 (dd, 1H, J 8.5, 2.0 Hz), 7.14 (d,
25 1H, J 3.0 Hz), 7.06 (dd, 1H, J 8.8, 3.0 Hz), 6.97 (d, 1H, J 8.8 Hz), 5.77 (dd, 1H, J 13.5, 2.7 Hz), 3.18 (dd, 1H, J -16.9, 13.5 Hz), 2.78 (dd, 1H, J -16.9, 2.7 Hz).

b) 2-(2,4-Dichlorophenyl)chroman-4,6-diol

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2-(2,4-Dichlorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.2 g of 2-(2,4-dichloro-phenyl)-6-hydroxychroman-4-one. The product was purified by column chromatography using heptane - ethyl acetate (2:1) as an eluant. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.89 (s, 1H), 7.66 (d, 1H, J 2.1 Hz), 7.64 (d, 1H, J 8.5 Hz), 7.51 (dd,
35 1H, J 2.1, 8.5 Hz), 6.89 (d, 1H, J 2.7 Hz), 6.63 (d, 1H, J 8.7 Hz), 6.56 (dd, 1H, J 2.7,

8.7 Hz), 5.50 (d, 1H, J 6.8 Hz), 5.37 (d, 1H, J 10.4 Hz), 4.90 (m, 1H), 2.32 (m, 1H), 1.80 (m, 1H).

c) 2-(2,4-Dichlorophenyl)chroman-6-ol

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2-(2,4-Dichlorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 625 mg of 2-(2,4-dichlorophenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 7.65 (d, 1H, J 2.2 Hz), 7.57 (d, 1H, J 8.4 Hz), 7.49 (dd, 1H, J 8.4, 2.2 Hz), 6.67-6.51 (m, 3H), 5.21 (dd, 1H, J 10.3, 2.1 Hz), 2.91 (m, 1H), 2.69 (m, 1H), 2.16 (m, 1H), 1.85 (m, 1H).

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d) 2-[2-(2,4-Dichlorophenyl)chroman-6-yloxy]-5-nitropyridine

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2-[2-(2,4-dichlorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 530 mg of 2-(2,4-dichlorophenyl)chroman-6-ol. The product was purified on preparative TLC-plate covered with silica gel using heptane - ethyl acetate (3:1) as an eluant. ¹H NMR (400 MHz, CDCl₃) δ: 9.06 (d, 1H, J 2.7 Hz), 8.47 (dd, 1H, J 9.0, 2.7 Hz), 7.56 (d, 1H, J 8.4 Hz), 7.41 (d, 1H, J 2.0 Hz), 7.33 (dd, 1H, J 8.4, 2.0 Hz), 7.02 (d, 1H, J 9.0 Hz), 6.99-6.92 (m, 3H), 5.39 (dd, 1H, J 10.4, 2.2 Hz), 3.06 (ddd, 1H, J -16.9, 11.9, 6.0 Hz), 2.83 (ddd, 1H, J -16.9, 5.3, 2.7 Hz), 2.34 (m, 1H), 1.89 (m, 1H).

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Example 12:

2-[2-(3-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(3-Chlorophenyl)-6-hydroxychroman-4-one

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2-(3-Chlorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 2.0 g of 2',5'-dihydroxyacetophenone and 1.85 g of 3-chlorobenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.47 (s, 1H), 7.62 (s, 1H), 7.51-7.45 (m, 3H), 7.12 (d, 1H, J 3.0 Hz), 7.05 (dd, 1H, J 8.8, 3.0 Hz), 6.98 (d, 1H, J 8.8 Hz), 5.58 (dd, 1H, J 13.1, 2.9 Hz), 3.18 (dd, 1H, J -16.9, 13.1 Hz), 2.81 (dd, 1H, J -16.9, 2.9 Hz).

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b) 2-(3-Chlorophenyl)chroman-4,6-diol

2-(3-Chlorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 730 mg of 2-(3-chlorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 7.50 (d, 1H, J 1.7 Hz), 7.46-7.38 (m, 3H), 6.88 (d, 1H, J 2.5 Hz), 6.62 (d, 1H, J 8.6 Hz), 6.55 (dd, 1H, J 8.6, 2.5 Hz), 5.44 (d, 1H, J 6.6 Hz), 5.15 (dd, 1H, J 11.8, 1.4 Hz), 4.87 (m, 1H), 2.29 (m, 1H), 1.85 (m, 1H).

c) 2-(3-Chlorophenyl)chroman-6-ol

2-(3-Chlorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 635 mg of 2-(3-chlorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.79 (s, 1H), 7.48 (d, 1H, J 0.7 Hz), 7.42-7.37 (m, 3H), 6.71-6.49 (m, 3H), 5.04 (m, 1H), 2.91 (m, 1H), 2.65 (m, 1H), 2.12 (m, 1H), 1.93 (m, 1H).

d) 2-[2-(3-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(3-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 590 mg of 2-(3-chlorophenyl)chroman-6-ol. The product was recrystallised from a 3:1 mixture of 2-propanol and ethyl acetate. ¹H NMR (400 MHz, CDCl₃) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.0, 2.9 Hz), 7.53 (s, 1H), 7.46-7.42 (m, 3H), 7.20 (d, 1H, J 9.0 Hz), 7.00 (dd, 1H, J 8.7, 2.7 Hz), 6.97 (d, 1H, J 2.7 Hz), 6.94 (d, 1H, J 8.7 Hz), 5.18 (dd, 1H, J 10.2, 2.2 Hz), 2.97 (ddd, 1H, J -17.0, 11.5, 5.9 Hz), 2.83 (ddd, 1H, J -17.0, 8.1, 4.5 Hz), 2.21 (m, 1H), 2.00 (m, 1H).

Example 13:

2-[2-(3,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(3,5-Difluorophenyl)-6-hydroxychroman-4-one

2-(3,5-Difluorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 1.0 g

of 2',5'-dihydroxyacetophenone and 1.12 g of 3,5-difluorobenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.47 (s, 1H), 7.30-7.23 (m, 3H), 7.12 (d, 1H, J 2.9 Hz), 7.06 (dd, 1H, J 8.8, 2.9 Hz), 7.00 (d, 1H, J 8.8 Hz), 5.60 (dd, 1H, J 13.1, 2.8 Hz), 3.15 (dd, 1H, J -16.8, 13.1 Hz), 2.85 (dd, 1H, J -16.8, 2.8 Hz).

b) 2-(3,5-Difluorophenyl)chroman-4,6-diol

2-(3,5-Difluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 800 mg of 2-(3,5-difluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.87 (s, 1H), 7.21-7.17 (m, 3H), 6.88 (d, 1H, J 2.4 Hz), 6.64 (d, 1H, J 8.7 Hz), 6.55 (dd, 1H, J 2.4, 8.7 Hz), 5.47 (d, 1H, J 7.0 Hz), 5.17 (d, 1H, J 10.5 Hz), 4.86 (m, 1H), 2.32 (m, 1H), 1.85 (m, 1H).

c) 2-(3,5-Difluorophenyl)chroman-6-ol

2-(3,5-Difluorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 500 mg of 2-(3,5-difluorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.82 (s, 1H), 7.20-7.14 (m, 3H), 6.68 (d, 1H, J 8.6 Hz), 6.53 (d, 1H, J 2.9 Hz), 6.50 (dd, 1H, J 8.6, 2.9 Hz), 5.05 (dd, 1H, J 9.8, 2.2 Hz), 2.88 (ddd, 1H, J -16.7, 10.8, 5.9 Hz), 2.62 (ddd, 1H, J -16.7, 8.9, 5.0 Hz), 2.15 (m, 1H), 1.93 (m, 1H).

d) 2-[2-(3,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(3,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 340 mg of 2-(3,5-difluorophenyl)chroman-6-ol. The product was purified on preparative TLC-plate covered with silica gel using toluene - ethyl acetate as an eluant and then crystallised from 2-propanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 7.23-7.19 (m, 4H), 7.01-6.95 (m, 3H), 5.18 (dd, 1H, J 10.0, 2.1 Hz), 2.97 (ddd, 1H, J -16.9, 10.9, 5.7 Hz), 2.76 (ddd, 1H, J -16.9, 8.4, 4.7 Hz), 2.22 (m, 1H), 1.99 (m, 1H).

Example 14:**5 2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine****a) 2-(2,5-Difluorophenyl)-6-hydroxychroman-4-one**

2-(2,5-Difluorophenyl)-6-hydroxychroman-4-one was prepared as described
10 for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g
of 2',5'-dihydroxyacetophenone and 2.57 ml of 2,5-difluorobenzaldehyde. The
product was recrystallised from acetic acid. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.46
(s, 1H), 7.53 (m, 1H), 7.36-7.30 (m, 2H), 7.14 (d, 1H, J 3.0 Hz), 7.05 (dd, 1H, J 8.8,
3.0 Hz), 6.97 (d, 1H, J 8.8 Hz), 5.76 (dd, 1H, J 13.6, 2.7 Hz), 3.26 (dd, 1H, J -16.8,
15 13.6 Hz), 2.76 (dd, 1H, J -16.8, 2.7 Hz).

b) 2-(2,5-Difluorophenyl)chroman-4,6-diol

2-(2,5-Difluorophenyl)chroman-4,6-diol was prepared as described for 2-
20 phenylchroman-4,6-diol in Example 8(a) starting from 1.0 g of 2-(2,5-difluoro-
phenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.87 (s, 1H),
7.39-7.22 (m, 3H), 6.89 (d, 1H, J 2.8 Hz), 6.63 (d, 1H, J 8.7 Hz), 6.56 (dd, 1H, J 8.7,
2.8 Hz), 5.50 (d, 1H, J 6.8 Hz), 5.35 (d, 1H, J 11.2 Hz), 4.89 (m, 1H), 2.28 (m, 1H),
1.95 (m, 1H).

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c) 2-(2,5-Difluorophenyl)chroman-6-ol

2-(2,5-Difluorophenyl)chroman-6-ol was prepared as described for 2-(3-
fluorophenyl)chroman-6-ol in Example 9(c) starting from 420 mg of 2-(2,5-
30 difluorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.82 (s, 1H),
7.34-7.22 (m, 3H), 6.71-6.51 (m, 3H), 5.20 (m, 1H), 2.93 (m, 1H), 2.68 (m, 1H),
2.11 (m, 1H), 1.98 (m, 1H).

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d) 2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 100 mg of 2-(2,5-difluorophenyl)chroman-6-ol. The product was recrystallised from 2-propanol. ¹H NMR (400 MHz, CDCl₃) δ: 9.07 (dd, 1H, J 2.8, 0.4 Hz), 8.47 (dd, 1H, J 9.1, 2.8 Hz), 7.26 (m, 1H), 7.05-6.91 (m, 6H), 5.35 (dd, 1H, J 10.3, 1.5 Hz), 3.04 (ddd, 1H, J -16.9, 11.7, 6.0 Hz), 2.82 (ddd, 1H, J -16.9, 5.2, 3.0 Hz) 2.29 (m, 1H), 2.01 (m, 1H).

Example 15:

2-[2-(3-Bromophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(3-Bromophenyl)-6-hydroxychroman-4-one

2-(3-Bromophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 2.3 ml of 3-bromobenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.41 (s, 1H), 7.50 (m, 1H), 7.59-7.53 (m, 2H), 7.39 (m, 1H) 7.12 (d, 1H, J 2.9 Hz), 7.05 (dd, 1H, J 8.8, 2.9 Hz), 6.98 (d, 1H, J 8.8 Hz), 5.57 (dd, 1H, J 13.0, 2.9 Hz), 3.12 (dd, 1H, J -16.9, 13.0 Hz), 2.81 (dd, 1H, J -16.9, 2.9 Hz).

b) 2-(3-Bromo-phenyl)-chroman-4,6-diol

2-(3-Bromophenyl)-chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.0 g of 2-(3-bromophenyl)-6-hydroxychroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.83 (s, 1H), 7.63 (m, 1H) 7.53 (m, 1H) 7.46 (m, 1H) 7.37 (m, 1H), 6.88 (d, 1H, J 2.9 Hz), 6.62 (d, 1H, J 8.7 Hz), 6.55 (dd, 1H, J 8.7, 2.9 Hz), 5.42 (d, 1H, J 7.0 Hz), 5.14 (d, 1H, J 10.5 Hz), 4.86 (m, 1H), 2.29 (m, 1H), 1.84 (m, 1H).

c) 2-(3-Bromophenyl)chroman-6-ol

2-(3-Bromophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 700 mg of 2-(3-bromophenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.81 (s, 1H),

7.61 (m, 1H), 7.51 (m, 1H), 7.43 (m, 1H), 7.35 (m, 1H) 6.67-6.48 (m, 3H), 5.01 (m, 1H), 2.87 (m, 1H), 2.63 (m, 1H), 2.12 (m, 1H), 1.92 (m, 1H).

d) 2-[2-(3-Bromophenyl)chroman-6-yloxy]-5-nitropyridine

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2-[2-(3-Bromophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 339 mg of 2-(3-bromophenyl)chroman-6-ol. The product was filtered through silica gel using toluene - ethyl acetate as an eluant and then crystallised from 2-propanol. ¹H NMR (400 MHz, CDCl₃) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.2, 2.9 Hz), 7.66 (bs, 1H), 7.55 (m, 1H), 7.48 (m, 1H), 7.39 (m, 1H) 7.20 (d, 1H, J 9.2 Hz) 7.01-6.93 (m, 3H), 5.17 (dd, 1H, J 10.1, 2.2 Hz), 2.97 (m, 1H), 2.72 (m, 1H) 2.20 (m, 1H), 2.00 (m, 1H).

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Example 16:

2-[2-(4-Ethylphenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(4-Ethylphenyl)-6-hydroxychroman-4-one

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2-(4-Ethylphenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 1.0 g of 2',5'-dihydroxyacetophenone and 0.8 ml of 4-ethylbenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.43 (d, 2H, J 8.1 Hz), 7.25 (d, 2H, J 8.1 Hz), 7.11 (d, 1H, J 3.1 Hz), 7.03 (dd, 1H, J 8.9, 3.1 Hz), 6.93 (d, 1H, J 8.9 Hz), 5.51 (dd, 1H, J 13.0, 2.9 Hz), 3.15 (dd, 1H, J -16.9, 13.0 Hz), 2.75 (dd, 1H, J -16.9, 2.9 Hz), 2.62 (q, 2H, J 7.5 Hz), 1.18 (t, 3H, J 7.5 Hz).

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b) 2-(4-Ethylphenyl)chroman-4,6-diol

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2-(4-Ethylphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 474 mg of 2-(4-ethylphenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.81 (s, 1H), 7.34 (d, 2H, J 8.0 Hz) 7.22 (d, 2H, J 8.0 Hz), 6.88 (d, 1H, J 2.8 Hz), 6.57 (d, 1H, J 8.6 Hz), 6.53 (dd, 1H, J 8.6, 2.8 Hz), 5.39 (d, 1H, J 7.1 Hz), 5.06 (d, 1H, J 10.7 Hz), 4.86 (m, 1H), 2.61 (q, 2H, J 7.6 Hz), 2.29 (m, 1H), 1.84 (m, 1H), 1.19 (t, 3H, J 7.6 Hz).

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c) 2-(4-Ethylphenyl)chroman-6-ol

2-(4-Ethylphenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 425 mg of 2-(4-ethylphenyl)chroman-4,6-diol. The product was purified using heptane - ethyl acetate (3:1) as an eluant. ¹H NMR (400 MHz, CD₃OD) δ: 7.26 (d, 2H, J 8.2 Hz) 7.13 (d, 2H, J 8.2 Hz), 6.65 (d, 1H, J 8.6 Hz), 6.55 (dd, 1H, J 8.6, 2.8 Hz), 6.51 (d, 1H, J 2.8 Hz), 4.83 (dd, 1H, J 10.1, 2.3 Hz), 2.84 (m, 1H), 2.62 (m, 1H), 2.59 (q, 2H, J 7.6 Hz) 2.03 (m, 1H), 1.93 (m, 1H), 1.19 (t, 3H, J 7.6 Hz).

d) 2-[2-(4-Ethylphenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(4-Ethylphenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 205 mg of 2-(4-ethylphenyl)chroman-6-ol. The product was recrystallised from a mixture of 2-propanol and acetone. ¹H NMR (400 MHz, CDCl₃) δ: 9.04 (d, 1H, J 2.8 Hz), 8.60 (dd, 1H, J 9.1, 2.8 Hz), 7.36 (d, 2H, J 8.1 Hz) 7.24 (d, 2H, J 8.1 Hz), 7.20 (d, 1H, J 9.1 Hz), 7.00 (d, 1H, J 2.7 Hz) 6.96 (dd, 1H, J 8.8, 2.7 Hz), 6.89 (d, 1H, J 2.7 Hz), 5.11 (dd, 1H, J 10.1, 2.2 Hz), 2.98 (m, 1H), 2.75 (m, 1H), 2.62 (q, 2H, J 7.5 Hz) 2.16 (m, 1H), 2.01 (m, 1H), 1.19 (t, 3H, J 7.5 Hz).

Example 17:2-(3-Methyl-2-phenylchroman-6-yloxy)-5-nitropyridine

a) 6-Hydroxy-3-methyl-2-phenylchroman-4-one

6-Hydroxy-3-methyl-2-phenylchroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 2.0 g of 2,5-dihydroxypropiophenone and 1.63 ml of benzaldehyde. The product was purified by column chromatography using heptane - ethyl acetate (3:1) as an eluant. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.37 (s, 1H), 7.53 (m, 2H), 7.47-7.39 (m, 3H), 7.13 (d, 1H, J 3.1 Hz), 7.02 (dd, 1H, J 8.9, 3.1 Hz), 6.89(d, 1H, J 8.9 Hz), 5.17 (d, 1H, J 12.3), 3.18 (dq, 1H, J 12.3, 6.9 Hz), 0.84 (d, 3H, J 6.9 Hz).

b) 3-Methyl-2-phenylchroman-4,6-diol

3-Methyl-2-phenylchroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 474 mg of 6-hydroxy-3-methyl-2-phenylchroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.79 (s, 1H), 7.42-7.33 (m, 5H), 6.88 (bs, 1H), 6.53 (m, 2H), 5.37 (d, 1H, J 8.0 Hz), 4.70 (d, 1H, J 10.6 Hz), 1.94 (m, 1H), 0.73 (d, 3H, J 6.7 Hz).

c) 3-Methyl-2-phenylchroman-6-ol

3-Methyl-2-phenylchroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 605 mg of 3-methyl-2-phenylchroman-4,6-diol. ¹H NMR (400 MHz, CD₃OD) δ: 8.77 (s, 1H), 7.41-7.33 (m, 5H), 6.59-6.48 (m, 3H), 4.56 (d, 1H, J 9.2 Hz), 2.73 (dd, 1H, J -16.5, 5.0 Hz), 2.54 (dd, 1H, J -16.5, 5.8 Hz), 2.11 (m, 1H), 0.72 (d, 3H, J 6.6 Hz).

d) 2-(3-Methyl-2-phenylchroman-6-yloxy)-5-nitropyridine

2-(3-Methyl-2-phenylchroman-6-yloxy)-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 600 mg of 3-methyl-2-phenylchroman-6-ol. The product was purified by column chromatography using heptane - 2-propanol (20:1) as an eluant. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.8 Hz), 8.59 (dd, 1H, J 9.1, 2.8 Hz), 7.43-7.36 (m, 5H), 7.19 (d, 1H, J 9.1 Hz), 7.00 (d, 1H, J 2.6 Hz), 6.95 (dd, 1H, J 8.7, 2.6 Hz), 6.86 (d, 1H, J 8.7 Hz), 4.73 (d, 1H, J 9.3 Hz), 2.85 (dd, 1H, J -16.7, 5.0 Hz), 2.64 (dd, 1H, J -16.5, 10.9 Hz), 2.18 (m, 1H), 0.77 (d, 3H, J 6.7 Hz).

Example 18:

3-Methyl-6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-one

Methyl-6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-one was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 200 mg of 6-hydroxy-3-methyl-2-phenylchroman-4-one. The product was purified by column chromatography using heptane - ethyl acetate (2:1) as an eluant and then crystallised from a mixture of 2-propanol and acetone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.03 (d, 1H, J 2.9 Hz), 8.64 (dd, 1H, J 9.1, 2.9 Hz), 7.59-7.56 (m, 3H), 7.50-7.32 (m, 4H), 7.30 (d, 1H, J 9.1 Hz), 7.18 (d, 1H, J 8.9 Hz), 5.38 (d, 1H, J 12.5 Hz), 3.36 (dd, 1H, J 12.5, 6.9 Hz), 0.86 (d, 3H, J 6.9 Hz).

Example 19:**2-[2-(2-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine**

5 a) 2-(2-Fluorophenyl)-6-hydroxychroman-4-one

2-(2-Fluorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 2.0 g of 2',5'-dihydroxyacetophenone and 1.4 ml of 2-fluorobenzaldehyde. The product was
10 recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.45 (s, 1H), 7.67 (m, 1H), 7.47 (m, 1H), 7.32-7.25 (m, 2H), 7.14 (d, 1H, J 3.0 Hz), 7.04 (dd, 1H, J 8.9, 3.0 Hz), 6.95 (d, 1H, J 8.9 Hz), 5.77 (dd, 1H, J 13.5, 2.8 Hz), 3.26 (dd, 1H, J -16.9, 13.5 Hz), 2.76 (dd, 1H, J -16.9, 2.8 Hz).

15 b) 2-(2-Fluorophenyl)chroman-4,6-diol

2-(2-Fluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.19 g of 2-(2-fluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 7.56 (m,
20 1H), 7.40 (m, 1H), 7.28-7.21 (m, 2H), 6.89 (d, 1H, J 2.9 Hz), 6.60 (d, 1H, J 8.7 Hz), 6.54 (dd, 1H, J 8.7, 2.8 Hz), 5.46 (d, 1H, J 6.9 Hz), 5.35 (d, 1H, J 10.6 Hz), 4.89 (m, 1H), 2.26 (m, 1H), 1.98 (m, 1H).

25 c) 2-(2-Fluorophenyl)chroman-6-ol

2-(2-Fluorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 800 mg of 2-(2-fluorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 7.50 (m, 1H), 7.39 (m,
1H), 7.26-7.19 (m, 2H), 6.63 (m, 1H) 6.53-6.50 (m, 2H), 5.21 (dd, 1H, J, 10.2, 2.3
30 Hz), 2.98 (ddd, 1H, J -16.9, 11.2, 6.0 Hz), 2.66 (ddd, 1H, J -16.9, 5.0, 2.9 Hz) 2.11 (m, 1H), 1.99 (m, 1H).

d) 2-[2-(2-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine

35 2-[2-(2-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting

from 390 mg of 2-(2-fluorophenyl)chroman-6-ol. The product was purified by column chromatography using heptane- ethyl acetate (4:1) as an eluant. ¹H NMR (400 MHz, CDCl₃) δ: 9.04 (d, 1H, J 2.8 Hz), 8.60 (dd, 1H, J 9.1, 2.8 Hz), 7.56 (m, 1H), 7.43 (m, 1H), 7.30-7.22 (m, 2H), 7.20 (d, 1H, J 9.1 Hz), 7.02 (d, 1H, J 2.8 Hz), 6.98 (dd, 1H, J 8.7, 2.8 Hz), 6.91 (d, 1H, J 8.7 Hz), 5.37 (dd, 1H, J 10.4, 2.3 Hz), 3.04 (ddd, 1H, J -17.0, 11.5, 6.0 Hz), 2.82 (ddd, 1H, J -17.0, 5.1, 2.8 Hz) 2.18 (m, 1H), 2.08 (m, 1H).

Example 20:

2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-nitropyridine

a) 1-[2,5-Bis(benzyloxy)phenyl]ethanone

A mixture of 1-(2,5-dihydroxyphenyl)ethanone (3.16 g), benzyl chloride (7.04 g), potassium carbonate (12.4 g) and 18-Crown-6 (30 mg) in 2-butanone (50 ml) was heated under reflux for 5 hrs. After cooling the precipitate was filtered off. The filtrate was evaporated to dryness under reduced pressure and ether (50 ml) was added to it. The solution was washed twice with dilute sodium hydroxide solution, twice with dilute hydrochloric acid, dried over sodium sulphate and substantially evaporated to dryness under reduced pressure. The residue was triturated with cold n-heptane (30 ml), and the precipitate was filtered off with suction filtration giving after drying 2.85 g of 1-[2,5-Bis(benzyloxy)phenyl]ethanone. ¹H NMR (400 MHz, DMSO-d₆) δ = 2.50 (s, 3H), 5.08 (s, 2H), 5.18 (s, 2H), 7.20-7.50 (m, 13H).

b) Acetic acid 2,5-bis(benzyloxy)phenyl ester

A solution of 1-[2,5-bis(benzyloxy)phenyl]ethanone (2.25 g) and peracetic acid 40% (1.63 ml) in acetic acid (5.4 ml) was stirred at 60 °C for 1 h. After cooling to room temperature the precipitated product was collected by filtration, washed with cold ether and dried under reduced pressure. Acetic acid 2,5-bis(benzyloxy)phenyl ester was recrystallized from 2-propanol. Yield is 1.87 g. ¹H NMR (DMSO-d₆) δ = 2.23 (s, 1H), 5.03 (s, 2H), 5.05 (s, 2H), 6.84-7.44 (m, 13H).

c) 2,5-Bis(benzyloxy)phenol

A solution of acetic acid 2,5-bis(benzyloxy)phenyl ester (1.85 g) and 5M sodium hydroxide solution (10.6ml) in ethanol (11 ml) was heated under reflux for 6.5 hrs. After ethanol was evaporated under reduced pressure the clear solution was made acidic with diluted hydrochloric acid. The precipitated product was collected
5 by filtration, washed with cold water and dried under reduced pressure. Yield is 0.56 g. ¹H NMR (DMSO-d₆) δ = 4.97 (s, 2H), 5.01 (s, 2H), 6.34 (dd, J = 3.1, 8.8 Hz, 1H), 6.49(d, J = 3.1 Hz, 1H), 6.85 (d, J = 8.8 Hz, 1H), 7.28-7.46 (m, 10H), 9.1 (br s, 1H).

d) 2-[2,5-Bis(benzyloxy)phenoxy]-1-phenylethanone

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A mixture of 2,5-bis(benzyloxy)phenol (0.28 g), 2-bromoacetophenone (0.22 g), potassium hydrogen-carbonate (0.25 g) and 18-Crown-6 (3 mg) in acetonitrile (4.2 ml) was stirred at 22 °C for one week. The mixture was filtered and evaporated to dryness under reduced pressure. The residue was triturated with the mixture of
15 ether (8.2 ml) and water (1.4 ml) at the ice bath temperature. The product was collected by filtration, washed with cold ether and dried under reduced pressure. Yield is 0.14 g. ¹H NMR (DMSO-d₆) δ = 4.98 (s, 2H), 5.06 (s, 2H), 5.58 (s, 2H), 6.51 (dd, J = 8.9, 2.3 Hz, 1H), 6.68 (d, J = 2.3 Hz, 1H), 6.94 (d, J = 8.9 Hz, 1H), 7.28-8.03 (m, 15H).

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e) 2-[2,5-Bis(benzyloxy)phenoxy]-1-phenylethanol

To the solution of 2-[2,5-bis(benzyloxy)phenoxy]-1-phenylethanone (0.14 g) in methanol (0.5 ml) and tetrahydrofuran (1.9 ml) was added at the 0 °C temperature sodium borohydride (6.5 mg). The reaction was stirred 15 minutes at 0 °C and 2 hrs
25 at 22 °C temperature. After adding water (5 ml) methanol and tetrahydrofuran were evaporated off. After the residue was stirred at 22 °C 0.5 hr the product was filtered, washed with cold water and dried under reduced pressure. Yield is 0.09 g. ¹H NMR (DMSO-d₆) δ = 4.05 (m, 2H), 4.91 (m, 1H), 4.95 (s, 2H), 5.01 (s, 2H), 5.59 (d, J =
30 4.7 Hz, 1H), 6.47 (dd, J = 2.8, 8.8 Hz, 1H), 6.68 (d, J = 2.8 Hz, 1H), 6.89 (d, J = 8.8 Hz, 1H), 7.24-7.45 (m, 15H).

f) 2-(2-Hydroxy-2-phenylethoxy)benzene-1,4-diol

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A solution of 2-[2,5-bis(benzyloxy)phenoxy]-1-phenylethanol (3.9 g) in ethanol (175 ml) was hydrogenated in the presence of 10 % palladium on charcoal

(100 mg) at 30 psi. The catalyst was removed by filtration and the solvent was evaporated under reduced pressure. The residue was recrystallized from the mixture of toluene-ethyl acetate 8:1 (15 ml). The yield of 2-(2-Hydroxy-2-phenylethoxy)-benzene-1,4-diol is 1.2 g. ¹H NMR (DMSO-d₆) δ = 3.79 (dd, J = 9.6, 8.3 Hz, 1H), 4.00 (dd, J = 9.6, 3.6 Hz, 1H), 4.94 (ddd, J = 3.6, 8.3, 3.9 Hz, 1H), 5.66 (d, J = 3.9 Hz, 1H), 6.18 (dd, J = 8.5, 2.3 Hz, 1H), 6.34 (d, J = 2.3, 1H), 6.57 (d, J = 8.5, 1H), 7.26-7.47 (m, 5H), 7.97 (s, 1H), 8.66 (s, 1H).

g) 2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-ol

A solution of 2-(2-hydroxy-2-phenylethoxy)benzene-1,4-diol (1.2 g) in toluene (75 ml) was heated with Amberlyst 15 catalyst (0.5 g) under reflux for 7 hrs. After filtering the solvent was evaporated under reduced pressure. The residue was purified by column chromatography on silica gel (toluene/ethyl acetate/acetic acid = 8:1:1). The yield of 2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-ol is 0.5 g. ¹H NMR (DMSO-d₆) δ = 4.02 (dd, J = 8.5, 11.4 Hz, 1H), 4.35 (dd, J = 2.3, 11.4 Hz, 1H), 5.11 (dd, J = 8.5, 2.3 Hz, 1H), 6.29 (dd, J = 2.8, 8.5 Hz, 1H), 6.32 (d, J = 2.8 Hz, 1H), 6.75 (d, J = 8.5 Hz, 1H), 7.36-7.47 (m, 5H), 8.99 (s, 1H).

h) 2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-nitropyridine

A solution of 2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-ol (80 mg), 2-chloro-5-nitropyridine (56 mg) and potassium carbonate (52 mg) in dimethylformamide (1.0 ml) was stirred at 120 °C for 2 hrs. After cooling the mixture water (10 ml) was added and the precipitated product was filtered, washed with water and 2-propanol and dried under reduced pressure. Yield is 60 mg and mp 163-170 °C. ¹H NMR (DMSO-d₆) δ = 4.16 (dd, J = 8.5, 11.6 Hz, 1H), 4.47 (dd, J = 11.6, 2.6 Hz, 1H), 5.28 (dd, J = 2.6, 8.5 Hz, 1H), 6.75 (dd, J = 2.6, 8.8 Hz, 1H), 6.88 (d, J = 2.6 Hz, 1H), 7.05 (d, J = 8.8 Hz, 1H), 7.21 (d, J = 9.1 Hz, 1H), 7.39-7.52 (m, 5H), 8.60 (dd, J = 2.8, 9.1 Hz, 1H), 9.05 (d, J = 2.8 Hz, 1H).

Example 21:

2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-3-nitropyridine

2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-3-nitropyridine was prepared in the same way as 2-(2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-

nitropyridine above from 2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-ol (80 mg) and 2-chloro-3-nitropyridine (56 mg). Yield is 30 mg and mp <60 °C. ¹H NMR (DMSO-d₆) δ = 4.16 (dd, J = 8.6, 11.4 Hz, 1H), 4.46 (dd, J = 11.4, 2.5 Hz, 1H), 5.27 (dd, J = 2.5, 8.6 Hz, 1H), 6.73 (dd, J = 2.5, 8.6 Hz, 1H), 6.85 (d, J = 2.5 Hz, 1H), 7.03 (d, J = 8.6 Hz, 1H), 7.34-7.52 (m, 6H), 8.43 (dd, J = 1.9, 4.8 Hz, 1H), 8.55 (dd, J = 7.8, 1.9 Hz, 1H).

Example 22:

2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-trifluoromethyl-pyridine

2-(2,3-Dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-trifluoromethyl-pyridine was prepared in the same way as 2-(2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-yloxy)-5-nitropyridine above from 2,3-dihydro-2-phenyl-benzo[1,4]dioxin-6-ol (80 mg) and 2-chloro-5-(trifluoromethyl)pyridine (64 mg). Yield is 50 mg and mp 104-110 °C. ¹H NMR (DMSO-d₆) δ = 4.15 (dd, J = 8.3, 11.4 Hz, 1H), 4.46 (dd, J = 2.3, 11.4 Hz, 1H), 5.27 (dd, J = 2.3, 8.3 Hz, 1H), 6.72 (dd, J = 2.8, 8.8 Hz, 1H), 6.84 (d, J = 2.8 Hz, 1H), 7.03 (d, J = 8.8 Hz, 1H), 7.19 (d, J = 8.8 Hz, 1H), 7.39-7.52 (m, 5H), 8.20 (dd, J = 8.8, 2.6 Hz, 1H), 8.58 (d, J = 2.6 Hz, 1H).

Example 23:

5-Nitro-2-(6-phenyl-5,6,7,8-tetrahydro-naphthalen-2-yloxy)-pyridine

a) 6-Methoxy-2-phenyl-3,4-dihydro-2H-naphthalen-1-one

A mixture of palladium(II) acetate (0.57 g), rac-2,2'-bis(diphenylphosphino)-1,1'-binaphthyl (1.91 g) and potassium *tert*-butoxide (4.15 g) in dry toluene was stirred under argon for 10 minutes. Bromobenzene (5.34 g) and 6-methoxy-1-tetralone (3.0 g) solvated in dry toluene were added and the mixture was stirred at 100 °C for 2 h. The reaction mixture was cooled to room temperature and poured into saturated aqueous ammonium chloride and extracted with ethyl ether. Organic extract was washed with brine, dried and evaporated. The crude product was purified by flash chromatography on silica gel using toluene and toluene-ethyl acetate (9:1) as an eluant. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.87 (d, 1H, J 7.8 Hz), 7.16-7.33 (m, 5H), 6.91-6.94 (m, 2H), 3.85 (s, 3H), 3.82-3.88 (m, 1H), 3.06-3.14 (m, 1H), 2.92-2.98 (m, 1H), 2.23-2.38 (m, 2H).

b) 6-Hydroxy-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one

6-Methoxy-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one (1.0 g) was refluxed
5 with 47 % HBr (20 ml) until disappearance of the starting material. The mixture was
poured into water and extracted with ethyl acetate. Ethyl acetate was dried and
evaporated. The product was recrystallised from toluene. ¹H NMR (400 MHz, d₆-
DMSO) δ: 10.35 (s, 1H), 7.79 (d, 1H, J 8.6 Hz), 7.15-7.33 (m, 5H), 6.75 (dd, 1H, J
8.6, 2.4 Hz), 6.68 (d, 1H, J 2.3 Hz), 3.79-3.85 (m, 1H), 2.99-3.06 (m, 1H), 2.83-2.90
10 (m, 1H), 2.19-2.33 (m, 2H).

c) 6-Phenyl-5,6,7,8-tetrahydro-naphthalen-2-ol

To a solution of 6-hydroxy-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one (50
15 mg) in trifluoroacetic acid was added triethylsilane (98 mg). The mixture was heated
at 60 °C for 3 h. Solvent was evaporated, water added to the residue and the mixture
extracted with ethyl acetate. Organic extract was dried and evaporated. ¹H NMR (400
MHz, d₆-DMSO) δ: 9.02 (s, 1H), 7.18-7.32 (m, 5H), 6.87 (d, 1H, J 7.9), 6.50-6.53
(m, 2H), 2.68-2.92 (m, 5H), 1.94-1.99 (m, 1H), 1.81-1.89 (m, 1H).
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d) 5-Nitro-2-(6-phenyl-5,6,7,8-tetrahydro-naphthalen-2-yloxy)-pyridine

6-Phenyl-5,6,7,8-tetrahydro-naphthalen-2-ol (30 mg), 2-chloro-5-nitropyridi-
ne (21 mg) and potassium fluoride (23 mg) in dry dimethylformamide were heated at
25 120 °C until disappearance of the starting material. Water and 1 N HCl were added
and the mixture extracted with ethyl acetate. Ethyl acetate was washed with brine and
water, dried and evaporated. The product was recrystallised from toluene. ¹H NMR
(400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.4 Hz), 8.61 (dd, 1H, J 9.0, 2.5), 7.18-7.35
(m, 7H), 6.95-6.99 (m, 2H), 2.83-3.01 (m, 5H), 1.87-2.04 (m, 2H).
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Example 24:

6-(5-Nitro-pyridin-2-yloxy)-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one

6-(5-Nitro-pyridin-2-yloxy)-2-phenyl-3,4-dihydro-2*H*-naphthalen-1-one was
35 prepared as described for 5-nitro-2-(6-phenyl-5,6,7,8-tetrahydro-naphthalen-2-
yloxy)-pyridine in Example 23(d) using 50 mg 6-hydroxy-2-phenyl-3,4-dihydro-2*H*-

naphthalen-1-one, 33 mg 2-chloro-5-nitropyridine and 37 mg potassium fluoride. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.07 (d, 1H, J 2.8 Hz), 8.68 (dd, 1H, J 9.0, 2.9), 8.01 (d, 1H, J 8.5), 7.37 (d, 1H, J 9.1 Hz), 7.21-7.38 (m, 7H), 3.96-4.04 (m, 1H), 3.15-3.23 (m, 1H), 2.98-3.04 (m, 1H), 2.39-2.48 (m, 1H), 2.25-2.31 (m, 1H).

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Example 25:**3-N-Acetylamino-4-(2-phenylchroman-6-yloxy)-anisole**

3-Amino-4-(2-phenylchroman-6-yloxy)-anisole of Example 3 (0.174 g) was dissolved in 2 ml of dry pyridine under nitrogen. Acetic anhydride (0.15 ml) and 10 mol-% of 4-(N,N-dimethylamino)pyridine (DMAP) were added and the reaction solution was stirred for 2.5 hours at room temperature and quenched with addition of 1 ml of H₂O. Toluene was added into the reaction mixture and evaporated to dryness. Toluene evaporation was repeated. Product was purified by column chromatography (CH₂Cl₂ : EtOAc / 98 : 2). ¹H-NMR (400 MHz; d₆-DMSO): δ 9.38 (s, 1H), 7.68 (s, 1H), 7.47-7.25 (m, 5H), 6.85-6.66 (m, 4H), 6.63 (dd, 1H, J = 8.8 Hz, J = 3.0 Hz), 5.07 (dd, 1H, J = 9.9 Hz, J = 1.9 Hz), 3.71 (s, 3H), 3.0-2.85 (m, 1H), 2.75-2.62 (s, 1H), 2.20-2.10 (m, 1H), 2.05 (s, 3H), 2.08-1.90 (m, 1H).

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Example 26:**5-Amino-2-(2-phenylchroman-6-yloxy)-pyridine**

5-Nitro-2-(2-phenylchroman-6-yloxy)-pyridine of Example 1 (2.26g) was dissolved in 350 ml of glacial acetic acid. Zinc powder (8.48g) was added in few portions due to exothermic reaction. The mixture was stirred at room temperature for 2 hours and filtered. The zinc was washed with glacial acetic acid. The acid was evaporated and toluene was added and evaporated again. A product mixture was dissolved in CH₂Cl₂ and washed with 1M NaOH. Water phase was further washed with CH₂Cl₂. Both organic fractions were combined and dried over Na₂SO₄. Product was purified by column chromatography. ¹H-NMR (400 MHz; d₆-DMSO): δ 7.52 (d, 1H, J = 2.8 Hz), 7.46-7.30 (m, 5H), 7.05 (dd, 1H, J = 8.6 Hz, J = 3.0 Hz), 6.82-6.72 (m, 3H), 6.69 (d, 1H, J = 8.6 Hz), 5.08 (dd, 1H, J = 10.0 Hz; J = 2.1 Hz), 5.00 (s, 2H), 3.00-2.87 (m, 1H), 2.74-2.64 (m, 1H), 2.19-2.10 (m, 1H), 2.05-1.91 (m, 1H).

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Example 27:**5-N-Acetylamino-2-(2-phenylchroman-6-yloxy)-pyridine**

5-Amino-2-(2-phenylchroman-6-yloxy)-pyridine of Example 26 (0.955g) was dissolved in 8 ml of dry pyridine under nitrogen. DMAP (0.038g) was added. AcCl (0.26 ml) was added at room temperature into the reaction solution dropwise because of vigorous and exothermic reaction. The reaction was stirred for 90 minutes at room temperature and quenched with slow addition of 1ml of H₂O. 50 ml of toluene was added and evaporated to dryness. Toluene evaporation was repeated twice. Brownish product mixture was purified with column chromatography (EtOAc:CH₂Cl₂ / 60:40) to give of crystalline slightly yellowish product. The product was further purified with recrystallization from MeOH:H₂O (71:29). The precipitate was filtered and washed with 2 × 10 ml of ice cold MeOH:H₂O (1:1). ¹H-NMR (400 MHz; d₆-DMSO): δ 10.04 (s, 1H), 8.27 (d, 1H, *J* = 2.4 Hz), 8.01 (dd, 1H, *J* = 8.9 Hz; *J* = 2.8 Hz), 7.47-7.31 (m, 5H), 6.93 (d, 1H, *J* = 8.9 Hz), 6.85 (d, 2H, *J* = 6.8 Hz), 6.84 (s, 1H), 5.11 (dd, 1H, *J* = 10.1 Hz; *J* = 2.2 Hz), 3.02-2.90 (m, 1H), 2.71 (ddd, 1H, *J* = 16.8 Hz; *J* = 5.1 Hz, *J* = 1.8 Hz), 2.22-2.11 (m, 1H), 2.08-1.92 (m, 1H).

Example 28:**5-*N,N*-Diacetyl-amino-2-(2-phenylchroman-6-yloxy)-pyridine**

5-Amino-2-(2-phenylchroman-6-yloxy)-pyridine of Example 26 (0.40g) was dissolved in 1.5 ml of dry pyridine under nitrogen. DMAP (10 mol-%) and Ac₂O (1.0 ml) were added and the solution was stirred at room temperature for 2.5 hours. Toluene was added and evaporated to dryness. Toluene evaporation was repeated twice. Product was purified by column chromatography (EtOAc:CH₂Cl₂ / 60:40). ¹H-NMR (400 MHz; d₆-DMSO): δ 8.05 (d, 1H, *J* = 2.6 Hz), 7.78 (dd, 1H, *J* = 8.7 Hz; *J* = 2.6 Hz), 7.48-7.31 (m, 5H), 7.07 (d, 1H, *J* = 8.9 Hz), 6.99-6.85 (m, 3H), 5.13 (dd, 1H, *J* = 10.1 Hz; *J* = 1.9 Hz), 3.05-2.92 (m, 1H), 2.78-2.70 (m, 1H), 2.21 (s, 6H), 2.25-2.12 (m, 1H), 2.08-1.94 (m, 1H).

Example 29:**2-(2-Phenylchroman-6-yloxy)-aniline****a) 2-Nitro-1-(2-phenylchroman-6-yloxy)-benzene**

6-Hydroxyflavane (0.150 g) was dissolved in dry DMSO (5 ml) under nitrogen. KI (0.166 g) and KOH (0.074g) were added and solution was stirred at

room temperature for 15 minutes. 2-Chloro-1-nitrobenzene (0.104 g) was added and the solution was stirred at +90°C for 2.5 hours. Cooled solution was taken up with CH₂Cl₂ and washed with H₂O and 2M NaOH. Phases were separated and water phase was washed CH₂Cl₂. All organic phases were combined and washed with 1M HCl and then H₂O (until pH~7) and brine. Solution was dried over Na₂SO₄ and purified with filtration through the small silica column in CH₂Cl₂:*n*-heptane (60:40).
¹H-NMR (300 MHz; d₆-DMSO): δ 8.01 (dd, 1H, *J* = 8.1 Hz; *J* = 1.7 Hz), 7.69 (m, 1H), 7.50-7.31 (m, 5H), 7.31-7.24 (m, 1H), 7.04 (dd, 1H, *J* = 8.5 Hz; *J* = 1.2 Hz), 6.92 (s, 1H), 6.90 (s, 2H), 5.12 (dd, 1H, *J* = 10.1 Hz; *J* = 2.3 Hz), 2.95 (ddd, 1H, *J* = 16.9 Hz; *J* = 11.2 Hz, *J* = 5.8 Hz), 2.73 (ddd, 1H, *J* = 16.9 Hz; *J* = 3.2 Hz, *J* = 1.7 Hz), 2.25-2.10 (m, 1H), 2.08-1.90 (m, 1H).

b) 2-(2-Phenylchroman-6-yloxy)-aniline

2-Nitro-1-(2-phenylchroman-6-yloxy)-benzene (0.160 g) was mostly dissolved in 30 ml of glacial acetic acid. Zinc powder (1.190 g) was added in few portions and the mixture was stirred for 90 minutes at room temperature. The zinc was filtered and washed with glacial acetic acid and evaporated. The evaporation residue was taken up with toluene and evaporated again. The residue was dissolved in CH₂Cl₂ and washed with 1M NaOH. NaOH solution was further washed with CH₂Cl₂. Both organic fractions were combined and dried over Na₂SO₄. The purification of the crude product was done by elution in CH₂Cl₂ through a small silica column. ¹H-NMR (400 MHz; d₆-DMSO): δ 7.45-7.28 (m, 5H), 6.89-6.83 (m, 1H), 6.80 (d, 1H, *J* = 8.5 Hz), 6.77 (dd, 1H, *J* = 8.0 Hz; *J* = 1.7 Hz), 6.73-6.67 (m, 3H), 6.54-6.48 (m, 1H), 5.06 (dd, 1H, *J* = 10.1 Hz; *J* = 2.3 Hz), 4.85 (s, 2H), 2.99-2.87 (m, 1H), 2.73-2.61 (m, 1H), 2.19-2.09 (m, 1H), 2.03-1.90 (m, 1H).

Example 30:

5-Trifluoromethyl-2-(2-phenylchroman-6-yloxy)-aniline

a) 2-Nitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene

2-Nitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene was prepared as described for 2-nitro-1-(2-phenylchroman-6-yloxy)-benzene in Example 29(a) except that 6-hydroxyflavane (0.339 g) was used in 7 ml of dry DMSO under nitrogen. Also KI (0.374 g) and KOH (0.168g) and 4-chloro-3-nitro-1-trifluoro-

methylbenzene (0.24 ml) were added in similar manner. Product was purified by column chromatography (CH_2Cl_2 : *n*-heptane / 60:40). ^1H -NMR (300 MHz; d_6 -DMSO): δ 8.44 (d, 1H, $J = 2.1$ Hz), 7.99 (dd, 1H, $J = 9.0$ Hz; $J = 2.2$ Hz), 7.51-7.29 (m, 5H), 7.15 (d, 1H, $J = 8.7$ Hz), 7.09-6.91 (m, 3H), 5.15 (dd, 1H, $J = 10.1$ Hz; $J = 2.3$ Hz), 3.08-2.90 (m, 1H), 2.83-2.68 (m, 1H), 2.25-2.11 (m, 1H), 2.09-1.91 (m, 1H).

b) 5-Trifluoromethyl-2-(2-phenylchroman-6-yloxy)-aniline

2-Nitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene (0.311 g) was reduced in 25 ml of glacial acetic acid with zinc (1.48 g) as described for 2-(2-phenylchroman-6-yloxy)-aniline in Example 29. Product was purified by column chromatography (CH_2Cl_2 : *n*-heptane / 70:30 as the eluant). ^1H -NMR (300 MHz; d_6 -DMSO): δ 7.48-7.28 (m, 5H), 7.06 (d, 1H, $J = 2.2$ Hz), 6.86 (dd, 1H, $J = 7.8$ Hz, $J = 1.5$ Hz), 6.85-6.56 (m, 3H), 6.72 (d, 1H, $J = 8.4$ Hz), 5.40 (s, 2H), 5.10 (dd, 1H, $J = 10.0$ Hz; $J = 2.3$ Hz), 3.04-2.87 (m, 1H), 2.78-2.65 (m, 1H), 2.24-2.10 (m, 1H), 2.08-1.89 (m, 1H).

Example 31:

5-Amino-2-(2-phenylchroman-6-yloxy)-aniline

a) 2,4-Dinitro-1-(2-phenylchroman-6-yloxy)-benzene

2,4-Dinitro-1-(2-phenylchroman-6-yloxy)-benzene was prepared as described for 2-nitro-1-(2-phenylchroman-6-yloxy)-benzene in Example 29(a) except that 6-hydroxyflavane (0.0226 g) was used in 5 ml of dry DMSO under nitrogen. Also KI (0.249 g) and KOH (0.112g) and 2,4-dinitrochlorobenzene (0.210 mg) were added in similar manner. Product was purified by column chromatography (CH_2Cl_2 : *n*-heptane / 75:25 as the eluant). ^1H -NMR (400 MHz; d_6 -DMSO): δ 8.88 (d, 1H, $J = 2.8$ Hz), 8.45 (dd, 1H, $J = 9.4$ Hz, $J = 2.9$ Hz), 7.48-7.30 (m, 5H), 7.14 (d, 1H, $J = 9.3$ Hz), 7.10 (d, 1H, $J = 2.8$ Hz), 7.05 (dd, 1H, $J = 8.8$ Hz, $J = 2.9$ Hz), 6.98 (d, 1H, $J = 8.7$ Hz), 5.16 (dd, 1H, $J = 10.2$ Hz, $J = 2.1$ Hz), 3.08-2.93 (m, 1H), 2.83-2.71 (m, 1H), 2.25-2.13 (m, 1H), 2.08-1.94 (m, 1H).

b) 5-Amino-2-(2-phenylchroman-6-yloxy)-aniline

2,4-Dinitro-1-(2-phenylchroman-6-yloxy)-benzene (0.04 g) was dissolved in 12 ml of glacial acetic acid and zinc (0.131 g) was added. The reaction stirred at room temperature for 30 minutes. Workup was done as described for 2-(2-phenylchroman-6-yloxy)-aniline in Example 29. Product was purified by column chromatography (CH₂Cl₂ : Et₃N / 96:4 as the eluant). ¹H-NMR (300 MHz; d₆-DMSO): δ 7.48-7.28 (m, 5H), 6.74 (d, 1H, *J* = 8.8 Hz), 6.63 (dd, 1H, *J* = 8.8 Hz, *J* = 2.9 Hz), 6.57 (d, 1H, *J* = 2.8 Hz), 6.49 (d, 1H, *J* = 8.4 Hz), 6.02 (d, 1H, *J* = 2.6 Hz), 5.81 (dd, 1H, *J* = 8.4 Hz, *J* = 2.6 Hz), 5.03 (dd, 1H, *J* = 10.0 Hz, *J* = 2.3 Hz), 4.62 (s, 2H), 4.48 (s, 2H), 2.99-2.82 (m, 1H), 2.71-2.57 (m, 1H), 2.20-2.06 (m, 1H), 2.03-1.86 (m, 1H)

Example 32:

5-Cyano-2-(2-phenylchroman-6-yloxy)-aniline

a) 4-Cyano-2-nitro-1-(2-phenylchroman-6-yloxy)-benzene

4-Cyano-2-nitro-1-(2-phenylchroman-6-yloxy)-benzene was prepared as described for 2-nitro-1-(2-phenylchroman-6-yloxy)-benzene in Example 29(a) except that 6-hydroxyflavane (0.453 g) was used in 10 ml of dry DMSO under nitrogen. Also KI (0.498 g) and KOH (0.224 g) and 4-chloro-3-nitro benzonitrile (0.365 mg) were added in similar manner. Product was purified by column chromatography (CH₂Cl₂ : *n*-heptane / 90:10 as the eluant). ¹H-NMR (400 MHz; d₆-DMSO): δ 8.64 (d, 1H, *J* = 2.0 Hz), 8.06 (dd, 1H, *J* = 8.8 Hz, *J* = 2.1 Hz), 7.08 (d, 1H, *J* = 8.8 Hz), 7.07-6.93 (m, 3H), 5.15 (dd, 1H, *J* = 10.1 Hz, *J* = 2.1 Hz), 3.05-2.91 (m, 1H), 2.82-2.70 (m, 1H), 2.24-2.12 (m, 1H), 2.08-1.92 (m, 1H).

b) 5-Cyano-2-(2-phenylchroman-6-yloxy)-aniline

4-Cyano-2-nitro-1-(2-phenylchroman-6-yloxy)-benzene (0.155 g; 0.4 mmol) was reduced to the corresponding aniline as described for 2-(2-phenylchroman-6-yloxy)-aniline in Example 29 except that 40 ml of glacial acetic acid and 0.93 g of zinc powder were used. Product was purified by column chromatography (100% CH₂Cl₂ as the eluant). ¹H-NMR (300 MHz; d₆-DMSO): δ 7.57-7.28 (m, 5H), 7.06 (d, 1H, *J* = 2.0 Hz), 6.95-6.78 (m, 4H), 6.65 (d, 1H, *J* = 8.3 Hz), 5.46 (s, 2H), 5.11 (dd, 1H, *J* = 10.0 Hz, *J* = 2.1 Hz), 3.03-2.88 (m, 1H), 2.78-2.66 (m, 1H), 2.23-2.10 (m, 1H), 2.08-1.89 (m, 1H).

Example 33:**N-Acetyl-2-(2-Penylchroman-6-yloxy)-aniline**

2-(2-Penylchroman-6-yloxy)-aniline (0.093 g) was dissolved in 1 ml of dry
5 pyridine under nitrogen. DMAP (10 mol-%) and acetic acid anhydride (0.1 ml) were
added and the solution was stirred for 4 hours at room temperature followed with
quenching with 0.5 ml of H₂O. The solution was evaporated to dryness and toluene
was added and evaporated again. Toluene evaporation was repeated. Product was
10 purified by column chromatography (CH₂Cl₂ : *i*-PrOH / 98:2 as the eluant) and
recrystallized from 0.5 ml of heated absolute ethanol by cooling and adding 0.5 ml
of. ¹H-NMR (400 MHz; d₆-DMSO): δ 9.43 (s, 1H), 7.96 (m, 1H), 7.48-7.30 (m, 5H),
7.08-6.99 (m, 2H), 6.89-6.74 (m, 4H), 5.10 (dd, 1H, *J* = 9.9, *J* = 2.0), 3.03-2.88 (m,
1H), 2.76-2.65 (m, 1H), 2.21-2.11 (m, 1H), 2.10-1.91 (m, 1H), 2.06 (s, 3H).

Example 34:**3-Nitro-2-(2-phenylchroman-6-yloxy)-pyridine**

6-Hydroxyflavane (0.150 g) was dissolved in 3 ml of dry DMF under
nitrogen. KF (0.117 g) was added and the solution was stirred for 30 minutes at
20 +120°C. The solution was cooled a bit and 2-chloro-3-nitropyridine was added
(0.212 g) and stirred for 7 hours at +120°C and overnight at room temperature. The
reaction mixture was taken up with EtOAc and 1M HCl and water were added and
phases separated. Organic phase was washed with water and pH was adjusted to 7
with 1M NaOH. Organic phase was washed with water, brine and dried over Na₂SO₄.
25 Product was purified by column chromatography (CH₂Cl₂ : *n*-heptane / 80:20).
¹H-NMR (400 MHz; d₆-DMSO): δ 8.55 (dd, 1H, *J* = 7.9 Hz, *J* = 1.7 Hz), 8.42 (dd,
1H, *J* = 4.9 Hz, *J* = 1.7 Hz), 7.51-7.29 (m, 6H), 7.02-6.92 (m, 2H), 6.88 (d, 1H, *J* =
8.7 Hz), 5.14 (dd, 1H, *J* = 10.0 Hz, *J* = 2.1 Hz), 3.05-2.92 (m, 1H), 2.78-2.68 (m,
1H), 2.22-2.13 (m, 1H) 2.07-1.95 (m, 1H).

30

Example 35:**3-Amino-5-(trifluoromethyl)-2-(2-phenylchroman-6-yloxy)-aniline**

a) 2,6-Dinitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene

35

2,6-Dinitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene was prepared as described for 2-nitro-1-(2-phenylchroman-6-yloxy)-benzene in Example 29(a) except that 6-hydroxyflavane (0.453 g) was used in 10 ml of dry DMSO under nitrogen. Also KI (0.498 g) and KOH (0.224 g) and 4-chloro-3,5-dinitro benzotri-
5 fluoride (0.541 mg) were added in similar manner. ¹H-NMR (400 MHz; d₆-DMSO): δ 8.87 (s, 2H), 7.46-7.29 (m, 5H), 6.89-6.75 (m, 4H), 5.09 (dd, 1H, *J* = 10.3 Hz, *J* = 2.1 Hz), 2.98-2.85 (m, 1H), 2.75-2.62 (m, 1H), 2.19-2.08 (m, 1H), 2.03-1.88 (m, 1H).

b) 3-Amino-5-(trifluoromethyl)-2-(2-phenylchroman-6-yloxy)-aniline

10 2,6-Dinitro-1-(2-phenylchroman-6-yloxy)-4-trifluoromethylbenzene (0.198 g; 0.43 mmol) was reduced to corresponding diamino compound as described for 2-(2-phenylchroman-6-yloxy)-aniline in Example 29 except that 25 ml of glacial acetic acid and 1.525 g of metallic zinc powder were used. Product was purified by column
15 chromatography (100% CH₂Cl₂ as the eluant). ¹H-NMR (400 MHz; d₆-DMSO): δ 7.45-7.28 (m, 5H), 6.76 (d, 1H, *J* = 8.8 Hz), 6.65 (dd, 1H, *J* = 8.7 Hz, *J* = 3.0 Hz), 6.62 (d, 1H, *J* = 2.8 Hz), 5.04 (dd, 1H, *J* = 10.0 Hz, *J* = 2.1 Hz), 4.98 (s, 4H), 2.98-2.86 (m, 1H), 2.70-2.60 (m, 1H), 2.18-2.09 (m, 1H), 2.00-1.88 (m, 1H).

20 **Example 36:**

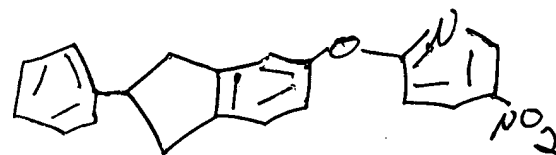
5-Succinimido-2-(2-phenylchroman-6-yloxy)-pyridine

5-Amino-2-(2-phenylchroman-6-yloxy)-pyridine of Example 26 (0.16 g) was dissolved in 7.5 ml of glacial acetic acid under nitrogen. Succinic anhydride (0.0563
25 g) was added and the solution refluxed 60 minutes and solution was cooled and evaporated to dryness. Toluene (25 ml) was added and evaporated again to dryness. Product was purified by column chromatography (CH₂Cl₂ : *i*-PrOH / 95:5 as the eluant). ¹H-NMR (400 MHz; d₆-DMSO): δ 8.03 (d, 1H, *J* = 2.6 Hz), 7.73 (dd, 1H, *J* = 8.7 Hz, *J* = 2.6 Hz), 7.48-7.31 (m, 5H), 7.10 (d, 1H, *J* = 8.7 Hz), 6.95 (d, 1H, *J* = 2.6
30 Hz), 6.92 (dd, 1H, *J* = 8.7 Hz, *J* = 2.7 Hz), 6.88 (d, 1H, *J* = 8.6 Hz), 5.13 (dd, 1H, *J* = 10.2 Hz, *J* = 2.2 Hz), 3.05-2.92 (m, 1H), 2.84-6.68 (m, 1H), 2.79 (s, 4H), 2.22-2.12 (m, 1H), 2.08-1.93 (m, 1H).

Example 37:

35 5-(*O,O'*-Diacetyl-L-tartaricimido)-2-(2-phenylchroman-6-yloxy)-pyridine

5-(*O,O'*-Diacetyl-L-tartaricimido)-2-(2-phenylchroman-6-yloxy)-pyridine was prepared as described for 5-succinimido-2-(2-phenylchroman-6-yloxy)-pyridine in Example 36 except that 5-amino-2-(2-phenylchroman-6-yloxy)-pyridine (0.318 g) and (+)-diacetyl-L-tartaric acid anhydride (0.227 g) were refluxed in 15 ml of glacial acetic acid for 60 minutes. ¹H-NMR (400 MHz; d₆-DMSO): δ 8.10 (d, 1H, *J* = 2.6 Hz), 7.79 (dd, 1H, *J* = 8.7 Hz, *J* = 2.6 Hz), 7.49-7.30 (m, 5H), 7.13 (d, 1H, *J* = 8.8 Hz), 6.97 (d, 1H, *J* = 2.5 Hz), 6.93 (dd, 1H, *J* = 8.7 Hz, *J* = 2.7 Hz), 6.88 (d, 1H, *J* = 8.7 Hz), 6.08 (s, 2H), 5.13 (dd, 1H, *J* = 10.1 Hz, *J* = 1.9 Hz), 3.06-2.92 (m, 1H), 2.80-2.69 (m, 1H), 2.23-2.12 (m, 1H), 2.18 (s, 6H), 2.08-1.93 (m, 1H).

Example 38:**5-Nitro-2-(2-phenylindan-5-yloxy) pyridine**

a) 3-(4-Methoxyphenyl)-2-phenylacrylic acid

Triethylamine was added to solution of p-anisaldehyde (10 g) and phenylacetic acid (10 g) in acetic anhydride (25 ml). Reaction mixture was stirred at 90°C for 8 h. Reaction mixture was cooled and water (600ml) solution of potassium carbonate (81 g) was added. After addition reaction mixture was heated at 60°C for an hour. Before neutralising with concentrated hydrochloric acid the reaction mixture was cooled below 10°C. Precipitate was filtered and washed with water. ¹H-NMR (400 MHz, d₆-DMSO): 12.6 (bs, 1H), 7.67 (s, 1H), 7.4-7.3 (m, 3H), 7.2-7.1 (m, 2H), 7.0-6.9 (m, 2H), 6.8-6.7 (m, 2H), 3.70 (s, 3H). (M)⁺ = 254 (100%).

b) 3-(4-Methoxyphenyl)-2-phenylpropionic acid

13 g of 3-(4-methoxyphenyl)-2-phenylacrylic acid was dissolved to 600 ml of ethyl acetate and 2.6 g of 10% palladium on charcoal was added under inert atmosphere. Starting material was hydrogenated at room temperature to give quantitative yield of 3-(4-methoxyphenyl)-2-phenylpropionic acid. ¹H-NMR (400 MHz, d₆-DMSO): 12.3 (bs, 1H), 7.32-7.20 (m, 5H), 7.1-7.0 (m, 2H), 6.8-6.7 (m, 2H), 3.79 (dd, 1H, *J* 6.9, 8.7 Hz), 3.70 (s, 3H), 3.22 (dd, 1H, *J* 8.7, 13.7 Hz), 2.87 (dd, 1H, *J* 6.9, 13.7 Hz).

c) 6-Methoxy-2-phenylindan-1-one

To solution of 3-(4-methoxyphenyl)-2-phenylpropionic acid (4.6 g) in dry methylenechloride (26 ml) was added two drops of dry DMF. Thionylchloride (3 ml) was added and reaction mixture was stirred at 40°C for 4 h. Solvent was evaporated under vacuum. Precipitate was dissolved to methylenechloride. Solution was cooled to 0-3°C. This solution and aluminium chloride (2.5g) were mixed slowly over 4 hours keeping temperature under 4°C. After mixing reaction mixture was stirred at room temperature for 2 h. Reaction was quenched by pouring to dilute ice cold hydrochloric acid. Layers were separated and water solution was extracted with methylenechloride. Combined organic layers were washed with water, dried and evaporated. Crude product was triturated to give 2.9 g of 6-Methoxy-2-phenylindan-1-one. ¹H-NMR (400 MHz, d₆-DMSO): 7.56 (d, 1H), 7.35-7.23 (m, 4H), 7.18-7.13 (m, 3H), 4.02 (dd, 1H, J 3.9, 8.0 Hz), 3.82 (s, 3H), 3.61 (dd, 1H, J 8.0, 17.2 Hz), 3.11 (dd, 1H, J 3.9, 17.2 Hz).

d) 5-Methoxy-2-phenylindane

5-Methoxy-2-phenylindane was prepared as described for 2-phenylchroman-6-ol in Example 1(a) using 600 mg of 6-methoxy-2-phenylindan-1-one. ¹H-NMR (400 MHz, d₆-DMSO): 7.32-7.27 (m, 4H), 7.21-7.18 (m, 1H), 7.13 (d, 1H, J 8.2 Hz), 6.83 (d, 1H, J 2.4 Hz), 6.72 (dd, 1H, J 2.4, 8.2 Hz), 3.72 (s, 3H), 3.64 (k, 1H, J 8.5 Hz), 3.23 (dt, 2H, J 8.5, 15.9 Hz), 2.92 (m, 2H).

e) 2-Phenylindan-5-ol

Mixture of 5-methoxy-2-phenylindane (200 mg) and concentrated HBr (4 ml) was refluxed for 5.5 h. Reaction mixture was allowed to cool to room temperature and 20 ml of ice water and it was extracted with methylenechloride. The combined organic layers were washed with brine and dried with Na₂SO₄. The solvents were evaporated to give 2-phenylindan-5-ol. ¹H-NMR (400 MHz, d₆-DMSO): 9.05 (bs, 1H), 7.3-7.28 (m, 4H), 7.26-7.15 (m, 1H), 7.0 (d, 1H, J 8.1 Hz), 6.64 (d, 1H, J 1.9 Hz), 6.55 (dd, 1H, J 1.9, 8.1 Hz), 3.60 (k, 1H, J 8.6 Hz), 3.18 (m, 2H), 2.86 (dt, 2H, J 8.6, 16 Hz).

f) 5-Nitro-2-(2-phenylindan-5-yloxy) pyridine

5-Nitro-2-(2-phenylindan-5-yloxy) pyridine was prepared as described for 2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 107 mg of 2-phenylindan-5-ol. ¹H-NMR (400 MHz, d₆-DMSO): 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 2.9, 9.1 Hz), 7.38-7.28 (m, 5H), 7.24-7.20 (m, 2H), 7.11 (d, 1H, J 2.2 Hz), 7.00 (dd, 1H, J 2.2, 8.0 Hz), 3.72 (k, 1H, J 8.9 Hz), 3.36-3.28 (m, 2H), 3.01 (dd, 2H, J 8.9, 15.3 Hz).

Example 39:**5-Methoxy-2-(2-phenylindan-5-yloxy)phenylamine**

10 a) 5-(4-Methoxy-2-nitrophenoxy)-2-phenylindane

5-(4-Methoxy-2-nitrophenoxy)-2-phenylindan was prepared as described for 6-(4-methoxy-2-nitrophenoxy)-2-phenylchroman in Example 3(a) using 575 mg of 2-phenylindan-5-ol. ¹H-NMR (400 MHz, d₆-DMSO): 7.60 (d, 1H, J 3.1 Hz), 7.35-7.28 (m, 5H), 7.25-7.16 (m, 3H), 6.86 (d, 1H, J 2.3 Hz), 6.78 (dd, 1H, J 2.3, 8.2 Hz), 3.84 (s, 3H), 3.67 (k, 1H, J 8.3 Hz), 3.25 (dd, 2H, J 8.3, 15.7 Hz), 2.95 (m, 2H). (M)⁺ = 361 (60%), 209 (100%)

20 b) 5-Methoxy-2-(2-phenylindan-5-yloxy)phenylamine

5-Methoxy-2-(2-phenylindan-5-yloxy)phenylamine was prepared as described for 5-methoxy-2-(2-phenylchroman-6-yloxy)phenylamine in Example 3(b) using 200 mg of 5-(4-methoxy-2-nitrophenoxy)-2-phenylindane. ¹H-NMR (400 MHz, d₆-DMSO): 7.33-7.27 (m, 4H), 7.23-7.18 (m, 2H), 6.82 (d, 1H, J 8.4 Hz), 6.81 (s, 1H), 6.77 (dd, 1H, J 2.3, 8.4 Hz), 6.69 (d, 1H, J 2.3 Hz), 6.48 (bd, 1H, J 6.4 Hz), 3.71 (s, 3H), 3.66 (k, 1H, J 8.3 Hz), 3.24 (dd, 2H, J 8.3, 15.6 Hz), 2.93 (m, 2H). (M)⁺ = 331 (100%)

Example 40:

30 **2-[2-(3-Fluorophenyl)-indan-5-yloxy]-5-methoxyphenylamine**

a) 2-(3-Fluorophenyl)indan-5-ol

2-(3-Fluorophenyl)indan-5-ol was prepared as described for 2-phenylindan-5-ol in Example 25(a-e) using 5 g of 3-fluorophenylacetic acid. ¹H-NMR (400 MHz, d₆-DMSO): 9.09 (s, 1H), 7.37-7.29 (m, 1H), 7.14-7.7.09 (m, 2H), 7.02-6.98 (m, 2H),

6.64 (d, 1H, J 1.7 Hz), 6.55 (dd, 1H, J 2.3, 8.1 Hz), 3.63 (k, 1H, J 8.3 Hz), 3.24-3.12 (m, 2H), 2.94-2.79 (m, 2H).

b) 5-Methoxy-2-[2-(3-fluorophenyl)indan-5-yloxy]phenylamine

5

5-Methoxy-2-[2-(3-fluorophenyl)indan-5-yloxy]phenylamine was prepared as described for 5-methoxy-2-(2-phenylchroman-6-yloxy)phenylamine in Example 3(a-b) using 500 mg of 2-(3-fluorophenyl)indan-5-ol. ¹H-NMR (400 MHz, CDCl₃): 7.26-7.21 (m, 1H), 7.16 (d, 1H, J 2.9 Hz), 7.09 (d, 1H, J 8.2 Hz), 7.02 (d, 1H, J 7.7 Hz),
10 6.96-6.88 (m, 4H), 6.83 (d, 1H, J 9.1 Hz), 6.72 (dd, 1H, J 2.9, 9.1 Hz), 3.69 (s, 3H), 3.65 (k, 1H, J 8.8 Hz), 3.26-3.19 (m, 2H), 2.94 (dd, 2H, J 8.8, 15.1 Hz).

Example 41:

2-(2-Phenylindan-5-yloxy)phenylamine

15

a) 5-(2-Nitrophenoxy)-2-phenylindane

5-(2-nitrophenoxy)-2-phenylindane was prepared as described for 6-(4-methoxy-2-nitrophenoxy)-2-phenylchroman in Example 3(a) using 200 mg 2-phenylindan-5-ol and 150 mg of 1-chloro-2-nitrobenzene. ¹H-NMR (400 MHz, d₆-DMSO): 8.04 (dd, 1H, J 1.6, 8.3 Hz), 7.68 (ddd, 1H, J 1.6, 7.4, 8.3 Hz), 7.36-7.27 (m, 6H), 7.24-7.17 (m, 1H), 7.11 (dd, 1H, J 1.1, 8.4 Hz), 6.98 (bd, 1H, J 2.3 Hz),
20 6.89 (dd, 1H, J 2.3, 8.1 Hz), 3.69 (k, 1H, J 8.5 Hz), 3.28 (dd, 2H J 8.5, 15.8 Hz), 3.05-2.95 (m, 2H).

25

b) 2-(2-Phenylindan-5-yloxy)phenylamine

2-(2-Phenylindan-5-yloxy)phenylamine was prepared as described for 2-(2-phenyl-chroman-6-yloxy)-phenylamine in Example 29(b) using 170 mg of 5-(2-nitrophenoxy)-2-phenylindane. ¹H-NMR (400 MHz, d₆-DMSO): 7.32-7.27 (m, 4H),
30 7.23-7.16 (m, 2H), 6.92-6.87 (m, 1H), 6.81-6.71 (m, 4H), 6.56-6.53 (m, 1H), 4.85 (bs, 2H), 3.65 (k, 1H, J 8.3 Hz), 3.23 (dd, 2H, J 8.3, 15.5 Hz), 2.97-2.88 (m, 2H).

Example 42:

35

2-(2-Phenylindan-5-yloxy)-5-trifluoromethylbenzene-1,3-diamine

a) 5-(2,6-Dinitro-4-trifluoromethylphenoxy)-2-phenylindan

Potassium-t-butoxide (117 mg) was added into a solution of 2-phenylindan-5-ol (200 mg) in dry DMF (3 ml). After stirring resulting mixture at room temperature for 30 min 4-chloro-3,5-dinitrobenzotrifluoride (275 mg) was added. Reaction mixture was stirred for a further three hours at 150°C. After cooling into room temperature water and ethyl acetate was added into the mixture. 1 M Hydrochloric acid was added into water layer and the solution was extracted with ethyl acetate. Combined organic layers were washed with water and brine, dried with Na₂SO₄ and evaporated. 5-(2,6-Dinitro-4-trifluoromethylphenoxy)-2-phenylindan was recrystallised from ethanol. ¹H-NMR (400 MHz, d₆-DMSO): 8.89 (s, 2H), 7.33-7.27 (m, 4H), 7.23-7.18 (m, 2H), 6.96 (d, 1H, J 2.5 Hz), 6.83 (dd, 1H, J 2.5, 8.2 Hz), 3.67 (k, 1H, J 8.6 Hz), 3.27-3.19 (m, 2H), 2.97-2.89 (m, 2H).

b) 2-(2-Phenylindan-5-yloxy)-5-trifluoromethylbenzene-1,3-diamine

2-(2-Phenylindan-5-yloxy)-5-trifluoromethylbenzene-1,3-diamine was prepared as described for 2-(2-phenylchroman-6-yloxy)-5-trifluoromethylbenzene-1,3-diamine in Example 35(b) using 230 mg of 5-(2,6-dinitro-4-trifluoromethylphenoxy)-2-phenylindan. ¹H-NMR (300 MHz, d₆-DMSO): 7.33-7.27 (m, 4H), 7.22-7.18 (m, 1H), 7.14 (d, 1H, J 8.1 Hz), 6.73-6.69 (m, 2H), 6.31 (s, 2H), 4.98 (s, 4H), 3.64 (k, 1H, J 8.5 Hz), 3.26-3.18 (m, 2H), 2.96-2.86 (m, 2H).

Example 43:25 6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ylamine

a) 6-Hydroxy-2-phenylchroman-4-one oxime

Sodium hydroxide (122 mg) was added into a cooled solution of 6-hydroxyflavanone (2 g) and hydroxylamine hydrochloride (900 mg) in ethanol (5 ml) and water (2 ml). Resulting mixture was refluxed for 6 hours and hydroxylamine hydrochloride (450 mg) and sodium hydroxide (61 mg) were added after every 45 min. After cooling into room temperature water (23 ml) and concentrated hydrochloric acid (5.6 ml) were added into the mixture. Precipitate was filtered, washed with water and dried in vacuum. ¹H-NMR (400 MHz, d₆-DMSO): 11.3 (s, 1H), 9.11 (s, 1H), 7.50-7.48 (m, 2H), 7.43-7.33 (m, 3H), 7.21 (d, 1H, J 2.9 Hz), 6.80 (d, 1H, J 8.8

Hz), 6.72 (dd, 1H, J 2.9, 8.8 Hz), 5.07 (dd, 1H, J 3.2, 11.9 Hz), 3.28 (dd, 1H, J 3.2, 17.1 Hz), 2.64 (dd, 1H, J 11.9, 17.1 Hz).

b) 4-Amino-2-phenylchroman-6-ol

5

A solution of 6-hydroxy-2-phenylchroman-4-one oxime (2.07 g) in 1,2-dimethoxy ethane (20 ml) was added into a cooled solution of titanium(IV)chloride (1.9 ml) and sodium borohydride (1.29 g) in 1,2-dimethoxy ethane (20 ml). Resulting mixture was stirred for a further 4 hours at room temperature. Reaction was
10 quenched with ice and pH was adjusted to 2-3. Resulting mixture was extracted with toluene. Water layer was made alkaline with solution of sodium hydroxide. Dark precipitate was filtered and washed with methanol. Methanol washings were evaporated to dryness, dissolved to water and neutralised and finally precipitate was filtered. Mixture of diastereomers of 4-amino-2-phenylchroman-6-ol was isolated as
15 its hydrochloride salt. (M)⁺ = 241 (4.4%), 224 (10%), 137 (100%).

c) 6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ylamine

6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-ylamine was prepared as
20 described for 2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 100 mg of hydrochloride salt of 4-amino-2-phenylchroman-6-ol. Mixture of diastereomers of 6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-ylamine was isolated as its hydrochloride salt. (M)⁺ = 363 (22%), 259 (100%), 242 (70%), 224 (64%), 223 (62%).

25

Example 44:

N-[6-(5-Nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-acetamide

Acetic anhydride (60 µl) was added into a cooled solution of 6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-ylamine hydrochloride (100 mg) and pyridine
30 (41 µl) in dry DMF. Resulting mixture was stirred for a further 20 hours at 0°C. Reaction was quenched with ice water and neutralised. Resulting mixture was extracted with dichloromethane, dried with Na₂SO₄ and evaporated. Mixture of diastereomers of *N*-[6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-acetamide was recrystallised from dichloromethane. (M)⁺ = 405 (24%), 301 (14%), 259 (100%),
35 242 (55%), 224 (96%).

Example 45:**Dimethyl-[6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-amine****a) 4-*N,N*-Dimethylamino-2-phenylchroman-6-ol**

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Sodium cyanoborohydride (274 mg) was added into a solution of free base of 4-amino-2-phenylchroman-6-ol (263 mg) and 37 % formaldehyde (1.4 ml) in acetonitrile (15 ml). After 30 min pH was adjusted to 6-7 with acetic acid. Resulting mixture was stirred at room temperature over night. Reaction mixture was evaporated to dryness, precipitate was dissolved to 10% solution of potassium hydroxide and extracted with methylene chloride. Combined organic layers were dried and evaporated to give 4-*N,N*-dimethylamino-2-phenylchroman-6-ol as mixture of diastereomers. ¹H-NMR (400 MHz, d₆-DMSO): 8.79 (bs, 1H, major and minor), 7.5-7.3 (m, 5H, major and minor), 6.96 (d, 1H, J 2.8 Hz, major), 6.75 (d, 1H, J 2.9 Hz, minor), 6.69 (d, 1H, 8.7 Hz, minor), 6.63-6.60 (m, 1H, major and minor), 6.53 (dd, 1H, J 2.8, 8.7 Hz, major), 5.17 (dd, 1H, J 2.7, 9.7 Hz, minor), 5.04 (d, 1H, J 10.7 Hz, major), 4.17 (dd, 1H, J 5.7, 11.4 Hz, major), 3.35 (m, 1H, minor), 2.24 (s, 3H, minor), 2.21 (s, 3H, major), 2.2-1.7 (m, 2H, major and minor).

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b) Dimethyl-[6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-amine

N,N-Dimethyl-[6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-amine was prepared as described for 2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 220 mg of 4-*N,N*-dimethylamino-2-phenylchroman-6-ol. *N,N*-Dimethyl-[6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl]-amine was isolated as a mixture of diastereomers. (M)⁺ = 391 (8%), 347 (8%), 346 (8%), 287 (68%), 147 (100%).

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Example 46:***N*-[6-(2-Phenylchroman-6-yloxy)pyridin-3-yl]methanesulfonamide**

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Pyridine (77 µl) and methanesufonyl chloride (32µl) were added into a cooled solution of 6-(2-phenylchroman-6-yloxy)pyridin-3-ylamine (121 mg) in dry THF (2 ml). After stirring resulting mixture at 0°C for additional 2 hours 1 M hydrochloric acid was added. Solution was extracted with ethyl acetate. Combined organic layers were dried with Na₂SO₄ and evaporated. *N*-[6-(2-Phenylchroman-6-yloxy)pyridin-3-yl]methanesulfonamide was recrystallised from diethyl ether. ¹H NMR (300 MHz,

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d_6 -DMSO) δ : 9.64 (s, 1H), 7.99 (d, 1H, J 2.8 Hz), 7.67 (dd, 1H, J 2.8, 8.8 Hz), 7.47-7.31 (m, 5H), 6.97 (d, 1H, J 8.8 Hz), 6.89-6.82 (m, 3H), 5.12 (dd, 1H J 2.2, 10.1 Hz), 3.0-2.9 (m, 1H), 2.98 (s, 3H), 2.77-2.69 (m, 1H), 2.20-2.13 (m, 1H), 2.04-1.96 (m, 1H).

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Example 47:1-Methyl-3-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl]thiourea

Solution of 6-(2-phenylchroman-6-yloxy)pyridin-3-ylamine (150 mg) and methyl isothiocyanate (94 μ l) in ethanol was refluxed for 10 hours. After cooling solvents were evaporated. Crude product of 1-methyl-3-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl]thiourea was purified by column chromatography (5% methanol in dichloromethane). ^1H NMR (400 MHz, d_6 -DMSO) δ : 9.45 (bs, 1H), 8.02 (d, 1H, J 2.7 Hz), 7.81 (dd, 1H, J 2.7, 8.8 Hz), 7.70 (bs, 1H), 7.47-7.38 (m, 4H), 7.36-7.32 (m, 1H), 6.94-6.86 (m, 4H), 5.12 (dd, 1H J 2.3, 10.1 Hz), 2.98-2.93 (m, 1H), 2.90 (d, 3H, J 4.3 Hz), 2.76-2.71 (m, 1H), 2.19-2.15 (m, 1H), 2.15-1.99 (m, 1H).

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Example 48:3-[6-(5-Nitropyridin-2-yloxy)chroman-2-yl]phenol

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a) 6-Hydroxy-2-(3-hydroxyphenyl)chroman-4-one

6-Hydroxy-2-(3-hydroxyphenyl)chroman-4-one was prepared as described for 6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was recrystallised from ethanol. ^1H NMR (400 MHz, d_6 -DMSO) δ : 9.50 (bs, 1H), 9.41 (bs, 1H), 7.22-7.17 (m, 1H), 7.11 (d, 1H, J 3.0 Hz), 7.03 (dd, 1H J 3.0, 8.9 Hz), 6.64 (d, 1H, J 8.9 Hz), 6.92-6.90 (m, 2H), 6.76-6.73 (m, 1H), 5.46 (dd, 1H J 2.9, 12.7 Hz), 3.09 (dd, 1H, J 12.7, 16.9 Hz), 2.75 (dd, 1H, J 2.9, 16.9 Hz).

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30 b) 2-(3-Hydroxyphenyl)chroman-4,6-diol

2-(3-Hydroxyphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(3-hydroxyphenyl)chroman-4-one. ^1H NMR (400 MHz, d_6 -DMSO) δ : 9.43 (bs, 1H), 8.88 (bs, 1H), 7.19-7.15 (m, 1H), 6.87 (d, 1H, J 2.7 Hz), 6.84-6.82 (m, 2H), 6.72-6.69 (m, 1H),

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6.58 (d, 1H, J 8.7 Hz), 6.53 (dd, 1H, J 2.7, 8.7), 5.01 (d, 1H, J 11.3 Hz), 4.86 (dd, 1H, J 6.2, 10.8 Hz), 2.25-2.19 (m, 1H), 1.88-1.75 (m, 1H).

c) 2-(3-Hydroxyphenyl)chroman-6-ol

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2-(3-Hydroxyphenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 2-(3-hydroxyphenyl)-chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.38 (s, 1H), 8.77 (s, 1H), 7.17-7.13 (m, 1H), 6.82-6.79 (m, 2H), 6.70-6.67 (m, 1H), 6.62 (d, 1H, J 8.6 Hz), 6.52-6.47 (m, 2H), 4.89 (dd, 1H, J 2.1, 9.9 Hz), 2.86-2.82 (m, 1H), 2.65-2.59 (m, 1H), 2.09-2.04 (m, 1H), 1.91-1.85 (m, 1H).

d) 3-[6-(5-Nitropyridin-2-yloxy)chroman-2-yl]phenol

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3-[6-(5-Nitropyridin-2-yloxy)chroman-2-yl]phenol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1 (b) starting from 2-(3-hydroxyphenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.44 (s, 1H), 9.04 (d, 1H, J 2.8 Hz), 8.60 (dd, 1H, J 2.8, 9.1 Hz), 7.21-7.16 (m, 2H), 7.00-6.94 (m, 2H), 6.91-6.84 (m, 3H), 6.73-6.70 (m, 1H), 5.06 (dd, 1H, J 2.1, 9.9 Hz), 2.99-2.92 (m, 1H), 2.75-2.69 (m, 1H), 2.17-2.01 (m, 1H), 2.00-1.93 (m, 1H).

Example 49:

6-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine

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6-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 830 mg of 2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine (Example 14(d)). ¹H NMR (300 MHz, d₆-DMSO) δ: 7.51 (d, 1H, J 2.9 Hz), 7.36-7.25 (m, 3H), 7.05 (dd, 1H, J 8.6, 2.9 Hz), 6.84-6.68 (m, 4H), 5.29 (d, 1H, J 8.6), 4.99 (s, 2H), 2.96 (m, 1H), 2.72 (m, 1H), 2.14 (m, 1H), 2.01 (m, 1H).

Example 50:

N-{6-[2-(2,5-Difluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

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N-{6-[2-(2,5-Difluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 370 mg of 6-[2-(2,5-difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 49). The product was purified on preparative TLC-plate covered with silica gel using ethyl acetate-heptane (4:1) as an eluant. ¹H NMR (400 MHz, CD₃OD) δ: 8.27 (d, 1H, J 2.7 Hz), 8.01 (dd, 1H, J 8.9, 2.7 Hz), 7.26 (m, 1H), 7.13 (m, 1H), 7.08 (m, 1H), 6.92-6.84 (m, 4H), 5.32 (dd, 1H, J 10.1, 1.6 Hz), 2.99 (ddd, 1H, J -16.9, 11.4, 5.9 Hz), 2.78 (ddd, 1H, J -16.9, 8.4, 5.1 Hz), 2.26 (m, 1H), 2.13 (s, 3H), 1.97 (m, 1H).

Example 51:

6-[2-(2-Fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine

6-[2-(2-Fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 240 mg of 2-[2-(2-fluorophenyl)chroman-6-yloxy]-5-nitropyridine (Example 19(d)). ¹H NMR (400 MHz, d₆-DMSO) δ: 7.52 (m, 1H), 7.51 (d, 1H, J 3.0 Hz), 7.41 (m, 1H), 7.28-7.24 (m, 2H), 7.05 (dd, 1H, J 8.6, 3.0 Hz), 6.81-6.73 (m, 3H), 6.70 (d, 1H, J 8.6 Hz), 5.31 (dd, 1H, J 10.3, 2.2 Hz), 5.00 (s, 2H), 2.98 (m, 1H), 2.72 (m, 1H), 2.15 (m, 1H), 2.06 (m, 1H).

Example 52:

N-{6-[2-(2-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

N-{6-[2-(2-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 220 mg of 6-[2-(2-fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 51). The product was recrystallised from a mixture of methanol and diethyl ether. ¹H NMR (400 MHz, d₆-DMSO) δ: 10.06 (s, 1H), 8.27 (d, 1H, J 2.7 Hz), 8.01 (dd, 1H, J 8.8, 2.7 Hz), 7.55 (m, 1H), 7.42 (m, 1H), 7.29-7.23 (m, 2H), 6.93 (d, 1H, J 8.8 Hz), 6.89-6.85 (m, 3H), 5.34 (dd, 1H, J 10.2, 2.2 Hz), 3.01 (m, 1H), 2.75 (m, 1H), 2.17 (m, 1H), 2.05 (m, 1H), 2.04 (s, 3H).

Example 53:

N-{6-[2-(2-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methanesulfonamide

N-{6-[2-(2-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methanesulfonamide was prepared as described for N-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl] methane sulfonamide in Example 46 starting from 400 mg of 6-[2-(2-fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 51). The product was crystallised from a mixture of methanol and diethyl ether. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.67 (s, 1H), 7.99 (d, 1H, J 2.8 Hz), 7.67 (dd, 1H, J 8.8, 2.8 Hz), 7.55 (m, 1H), 7.42 (m, 1H), 7.29-7.23 (m, 2H), 6.98 (dd, 1H, J 8.8 Hz), 6.92-6.84 (m, 3H), 5.35 (dd, 1H, J 10.4, 2.3 Hz), 3.01 (m, 1H), 2.99 (s, 3H), 2.76 (m, 1H), 2.16 (m, 1H), 2.04 (m, 1H).

Example 54:

6-[2-(3-fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine

6-[2-(3-fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 2,34 g of 2-[2-(3-fluorophenyl)chroman-6-yloxy]-5-nitropyridine (Example 9(d)). ¹H NMR (400 MHz, d₆-DMSO) δ: 7.51 (d, 1H, J 3.0 Hz), 7.44 (m, 1H), 7.30-7.25 (m, 2H), 7.16 (m, 1H), 7.05 (dd, 1H, J 8.6, 3.0 Hz), 6.83-6.73 (m, 3H), 6.69 (d, 1H, J 8.6 Hz), 5.13 (dd, 1H, J 10.0, 3.0 Hz), 5.00 (s, 2H), 2.93 (ddd, 1H, -16.8, 10.5, 5.3 Hz), 2.68 (ddd, 1H, J -16.8, 8.0, 4.4 Hz), 2.18 (m, 1H), 1.96 (m, 1H).

Example 55:

N-{6-[2-(3-fluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

N-{6-[2-(3-fluorophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 300 mg of 6-[2-(3-fluorophenyl)chroman-6-yloxy]pyridin-3-ylamine (Example 54). The product was recrystallised from a mixture of methanol and diethyl ether. ¹H NMR (400 MHz, d₆-DMSO) δ: 10.09 (s, 1H), 8.28 (d, 1H, J 2.7 Hz), 8.02 (dd, 1H, J 8.8, 2.7 Hz), 7.46 (m, 1H), 7.31-7.27 (m, 2H), 7.17 (m, 1H), 6.93 (d, 1H, J 8.8 Hz), 6.89-6.83 (m, 3H), 5.16 (dd, 1H, J 10.1, 2.1 Hz), 2.95 (ddd, 1H, J -16.5, 11.0, 6.5 Hz), 2.71 (ddd, 1H, J -16.5, 8.7, 4.4 Hz), 2.19 (m, 1H), 2.04 (s, 3H), 1.96 (m, 1H).

Example 56:

6-(5-Aminopyridin-2-yloxy)-2-phenylchroman-4-one

6-(5-Aminopyridin-2-yloxy)-2-phenylchroman-4-one was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 100 mg of 6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-one (Example 6). ¹H NMR (400 MHz, CD₃OD) δ: 7.62 (d, 1H, J 3.0 Hz), 7.51-7.49 (m, 2H), 7.42-7.33 (m, 3H), 7.25-7.18 (m, 3H), 7.06 (d, 1H, J 8.8 Hz), 6.76 (d, 1H, J 8.6 Hz), 5.50 (dd, 1H, J 13.0, 2.9 Hz), 3.08 (dd, 1H, J 17.0, 13.0 Hz), 2.82 (dd, 1H, J 17.0, 2.9 Hz).

Example 57:

Acetic acid 6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-yl ester

Acetanhydride (0.26 ml) was added dropwise into a solution of 100 mg of 6-(5-nitropyridin-2-yloxy)-2-phenylchroman-4-ol (Example 8(b)) in dry pyridine. The reaction mixture was refluxed for 1 ½ hours. It was then poured in ice-water and extracted with ethyl acetate. The organic phase was washed with 1 M HCl-solution, water and saturated NaCl-solution. It was then dried with MgSO₄ and evaporated to dryness. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.05 (d, 1H, J 2.8 Hz), 8.61 (dd, 1H, J 9.1, 2.8 Hz), 7.52-7.35 (m, 5H), 7.23 (d, 1H, J 9.1 Hz), 7.14-7.10 (m, 2H), 6.98 (d, 1H, J 8.5 Hz), 6.18 (dd, 1H, J 10.1, 6.4 Hz), 5.44 (dd, 1H, J 11.4, 1.4 Hz), 2.51 (m, 1H), 2.15 (m, 1H), 2.05 (s, 3H).

Example 58:

6-(2-Aminoethoxy)-2-phenylchroman-4-one methane sulfonate

a) 6-(2-Azidoethoxy)-2-phenylchroman-4-one

6-(2-Azidoethoxy)-2-phenylchroman-4-one was prepared as described for 6-(2-azidoethoxy)-2-phenylchroman in example 4(a) starting from 1.0 g of 6-hydroxyflavanone. ¹H NMR (400 MHz, CD₃OD) δ: 7.53-7.51 (m, 2H), 7.44-7.35 (m, 4H), 7.22 (dd, 1H, J 9.0, 3.1 Hz), 7.04 (d, 1H, J 9.0 Hz), 5.51 (dd, 1H, J 13.1, 3.0 Hz), 4.17 (t, 2H, J 4.9 Hz), 3.60 (t, 2H, J 4.9 Hz), 3.11 (dd, 1H, J 16.9, 13.1 Hz), 2.85 (dd, 1H, J 16.9, 3.0 Hz).

b) 6-(2-Aminoethoxy)-2-phenylchroman-4-one methane sulfonate

6-(2-Aminoethoxy)-2-phenylchroman-4-one methane sulfonate was prepared as described for 2-(2-phenylchroman-6-yloxy)ethylamine methane sulfonate in Example 4 (b) starting from 6-(2-azidoethoxy)-2-phenylchroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.91 (bs, 3H), 7.55-7.54 (m, 2H), 7.46-7.37 (m, 3H), 7.31 (d, 1H, J 3.1 Hz), 7.28 (dd, 1H, 8.8, 3.1 Hz), 7.12 (d, 1H, 8.8 Hz), 5.63 (dd, 1H, J 13.0, 3.0 Hz), 4.17 (t, 2H, J 5.1 Hz), 3.25 (dd, 1H, J -16.9, 13.0 Hz), 3.23 (m, 2H), 2.86 (dd, 1H, J -16.9, 3.0 Hz), 2.29 (s, 3H).

Example 59:

2-(3-Bromophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(3-Bromophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 215 mg of 2-(3-bromophenyl)chroman-4,6-diol (Example 15(b)). The product was recrystallised from a mixture of 2-propanol and acetone. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.8 Hz), 8.61 (dd, 1H, J 9.1, 2.8 Hz), 7.69 (m, 1H), 7.58-7.50 (m, 2H), 7.40 (m, 1H), 7.25 (d, 1H, J 2.7 Hz), 7.22 (d, 1H, 9.1 Hz), 7.02 (dd, 1H, J 8.8, 2.7 Hz), 6.91 (d, 1H, J 8.8 Hz), 5.65 (d, 1H, J 6.4 Hz), 5.33 (d, 1H, J 10.8 Hz), 4.97 (m, 1H), 2.36 (m, 1H), 1.94 (m, 1H).

Example 60:

2-(2-Fluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(2-Fluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 315 mg of 2-(2-fluorophenyl)chroman-4,6-diol (Example 19(b)). The product was recrystallised from a mixture of 2-propanol and acetone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.61 (m, 1H), 7.43 (m, 1H), 7.31-7.24 (m, 3H), 7.23 (d, 1H, J 9.1 Hz), 7.02 (dd, 1H, J 8.7, 3.1 Hz), 6.89 (d, 1H, J 8.7 Hz), 5.69 (d, 1H, J 6.3 Hz), 5.55 (d, 1H, J 11.9 Hz), 5.00 (m, 1H), 2.33 (m, 1H), 2.07 (m, 1H).

Example 61:

2-(2,5-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(2,5-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 200 mg of 2-(2,5-difluorophenyl)chroman-4,6-diol (Example 14(b)).

The product was purified on preparative TLC-plate covered with silica gel using
5 toluene –ethyl acetate (4:1) as an eluant. ¹H NMR (400 MHz, CDCl₃) δ: 9.03 (d, 1H, J 2.8 Hz), 8.50 (dd, 1H, J 9.1, 2.8 Hz), 7.36-7.33 (m, 2H), 7.08-6.95 (m, 5H), 5.50 (d, 1H, J 11.1 Hz), 5.09 (m, 1H), 2.53 (m, 1H), 2.05 (m, 1H).

Example 62:

10 2-(3-Fluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(3-Fluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 890 mg of 2-(3-fluorophenyl)chroman-4,6-diol (Example 9(b)). The product
15 was purified by column chromatography using gradient elution with ethyl acetate – heptane (20 %-> 33 %) and then crystallised from a mixture of 2-propanol and acetone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.48 (m, 1H), 7.35-7.31 (m, 2H), 7.26-7.19 (m, 3H), 7.02 (dd, 1H, J 8.8, 2.9 Hz), 6.91 (d, 1H, J 8.8 Hz), 5.67 (d, 1H, J 6.4 Hz), 5.34 (d, 1H, J 10.4 Hz), 4.98
20 (m, 1H), 2.36 (m, 1H), 1.99 (m, 1H).

Example 63:

2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-ethoxyphenylamine
hydrochloride

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a) 1-Chloro-4-ethoxy-2-nitrobenzene

4-Chloro-3-nitrobenzene (5,0 g) was dissolved in acetone and ethyl iodide (2,5 ml) and potassium carbonate (4,4 g) were added. The reaction mixture was
30 stirred at 40°C for 4 ½ hours. After cooling to room temperature the mixture was filtered and the filtrate was evaporated to dryness. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.64 (m, 2H), 7.27 (dd, 1H, J 9.0, 3.1 Hz), 4.12 (q, 2H, J 7.0 Hz), 1.34 (t, 3H, 7.0 Hz).

35 b) 2-(2,5-Difluorophenyl)-6-(4-ethoxy-2-nitrophenoxy)chroman

320 mg of 2-(2,5-Difluorophenyl)chroman-6-ol (Example 14(c)) was dissolved in dry DMF and potassium *tert*-butoxide (150 mg) was added. The resulting mixture was stirred for 30 minutes and 1-chloro-4-ethoxy-2-nitrobenzene (250 mg) was added. The reaction mixture was refluxed for 8 hours. After cooling to room temperature 1 M HCl-solution was added into the reaction mixture and it was extracted with ethyl acetate. The combined organic extracts were washed with water and saturated NaCl-solution. The product was purified by column chromatography using heptane - ethyl acetate (3:1) as an eluant. ¹H NMR (300 MHz, CDCl₃) δ: 7.43 (d, 1H, J 2.9Hz), 7.25 (m, 1H), 7.08-6.97 (m, 4H), 6.89-6.75 (m, 3H), 5.31 (d, 1H, J 9.1 Hz), 4.06 (q, 2H, J 7.0 Hz), 2.97 (m, 1H), 2.74 (m, 1H), 2.26 (m, 1H), 1.98 (m, 1H), 1.44 (t, 3H, 7.0 Hz).

c) 2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-ethoxyphenylamine hydrochloride

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2-[2-(2,5-Difluorophenyl)chroman-6-yloxy]-5-ethoxyphenylamine hydrochloride was prepared as described for 5-methoxy-2-(2-phenylchroman-6-yloxy)phenylamine hydrochloride in example 3(b) starting from 120 mg of 2-(2,5-difluorophenyl)-6-(4-ethoxy-2-nitrophenoxy)chroman. ¹H NMR (400 MHz, CDCl₃) δ: 7.26-7.22 (m, 2H), 7.01-6.74 (m, 7H), 5.22 (d, 1H, J 9.1 Hz), 3.90 (q, 2H, J 6.9 Hz), 2.90 (m, 1H), 2.68 (m, 1H), 2.19 (m, 1H), 1.90 (m, 1H), 1.34 (t, 3H, 6.9 Hz).

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Example 64:

5-Nitro-2-[2-(4-trifluoromethylphenyl)chroman-6-yloxy]pyridine

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a) 6-Hydroxy-2-(4-trifluoromethylphenyl)chroman-4-one

6-Hydroxy-2-(4-trifluoromethylphenyl)chroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 2,0 g of 2',5'-dihydroxyacetophenone and 2,1 ml of 4-trifluoromethylbenzaldehyde. The product was purified by column chromatography using heptane - ethyl acetate (2:1) as an eluant. Further purification was carried out by column chromatography using toluene-ethyl acetate (4:1) as an eluant. Finally the product was crystallised from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.47 (s, 1H), 7.82-7.76 (m, 4H), 7.13 (d, 1H, J 3.0 Hz), 7.06 (dd, 1H, J 8.8, 3.0 Hz), 6.99 (d, 1H, J 8.8 Hz),

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5.70 (dd, 1H, J 12.9, 2.9 Hz), 3.16 (dd, 1H, J -16.9, 12.9 Hz), 2.86 (dd, 1H, J -16.9, 2.9 Hz).

b) 2-(4-Trifluoromethylphenyl)chroman-4,6-diol

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2-(4-Trifluoromethylphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 860 mg of 2-(4-trifluoromethylphenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.86 (s, 1H), 7.77 (d, 2H, J 8.3 Hz), 7.68 (d, 2H, J 8.3 Hz), 6.89 (d, 1H, J 2.9 Hz), 6.63 (d, 1H, J 8.7 Hz), 6.56 (dd, 1H, J 8.7, 2.9 Hz), 5.45 (d, 1H, J 7.0 Hz), 5.26 (d, 1H, J 11.2 Hz), 4.90 (m, 1H), 2.32 (m, 1H), 1.85 (m, 1H).

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c) 2-(4-Trifluoromethylphenyl)chroman-6-ol

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2-(4-Trifluoromethyl phenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 730 mg of 2-(4-trifluoromethylphenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.82 (s, 1H), 7.75 (d, 2H, J 8.3 Hz), 7.65 (d, 2H, J 8.3 Hz), 6.67 (d, 1H, J 8.6 Hz), 6.53 (d, 1H, J 2.9 Hz) 6.51 (dd, 1H, 8.6, 2.9 Hz), 5.12 (d, 1H, J 8.3 Hz), 2.90 (m, 1H), 2.63 (m, 1H), 2.16 (m, 1H), 1.92 (m, 1H).

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d) 2-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 605 mg of 2-(4-trifluoromethylphenyl)chroman-6-ol. The product was purified column chromatography using 1,5 % ethyl acetate in toluene as an eluant and then crystallised from a mixture of 2-propanol and acetone. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 7.79 (d, 2H, J 8.2 Hz), 7.70 (d, 1H, J 8.2 Hz), 7.21 (d, 1H, J 9.1 Hz) 7.01 (dd, 1H, J 8.7, 2.7 Hz) 6.98 (d, 1H, J 2.7 Hz), 6.95 (d, 1H, 8.7 Hz), 5.29 (dd, 1H, J 10.1, 2.0 Hz), 3.00 (ddd, 1H, J -16.9, 10.1, 5.8 Hz), 2.4 (ddd, 1H, J -16.9, 8.4, 4.5 Hz) 2.24 (m, 1H), 1.99 (m, 1H).

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Example 65:

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6-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]pyridin-3-ylamine

6-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 275 mg of 2-[2-(4-trifluoromethylphenyl)chroman-6-yloxy]-5-nitropyridine (Example 64(d)). ¹H NMR (400 MHz, d₆-DMSO) δ: 7.78 (d, 2H, J 8.4 Hz), 7.68 (d, 2H, J 8.4 Hz), 7.52 (dd, 1H, J 2.9, 0.5 Hz), 7.06 (dd, 1H, 8.6, 2.9 Hz) 6.84 (m, 1H), 6.77-6.75 (m, 2H), 6.70 (dd, 1H, J 8.6, 0.5 Hz), 5.23 (dd, 1H, J 10.0, 2.0 Hz), 5.01 (s, 2H), 2.95 (ddd, 1H, -16.8, 11.1, 5.9 Hz), 2.69 (ddd, 1H, J -16.8, 8.5, 4.8 Hz), 2.21 (m, 1H), 1.97 (m, 1H).

Example 66:

N-{6-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

N-{6-[2-(4-Trifluoromethylphenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 140 mg of 6-[2-(4-trifluoromethylphenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 65). ¹H NMR (400 MHz, d₆-DMSO) δ: 10.11 (s, 1H), 8.29 (d, 1H, J 2.7 Hz), 8.02(dd, 1H, J 8.9, 2.7 Hz), 7.78 (d, 2H, J 8.3 Hz), 7.69 (d, 2H, J 8.3 Hz), 7.17 (m, 1H), 6.94-6.86 (m, 3H), 5.66 (d, 1H, J 8.2 Hz), 2.98 (m, 1H), 2.72 (m, 1H), 2.23 (m, 1H), 2.04 (s, 3H), 1.97 (m, 1H).

Example 67:

N-{6-[2-(3-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methane sulfonamide

N-{6-[2-(3-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methane sulfonamide was prepared as described for N-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl]methanesulfonamide in Example 46 starting from 300 mg of 6-[2-(3-fluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 54). The product was purified by passing through silica gel using ethyl acetate-heptane (5:1) as an eluant. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.67 (s, 1H), 7.99 (dd, 1H, J 2.8, 0.6 Hz), 7.67 (dd, 1H, J 8.8, 2.8 Hz), 7.46 (m, 1H), 7.31-7.27 (m, 2H), 7.17 (m, 1H), 6.98 (dd, 1H, J 8.8, 0.6 Hz) 6.90-6.88 (m, 3H), 5.16 (dd, 1H, J 10.0, 2.2 Hz), 2.99 (s, 3H), 2.96 (m, 1H), 2.72 (m, 1H), 2.20 (m, 1H), 1.99 (m, 1H).

Example 68:2-(4-Chlorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

a) 6-Hydroxy-2-(4-chlorophenyl)chroman-4-one

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6-Hydroxy-2-(4-chlorophenyl)chroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 2.8 g of 4-chlorobenzaldehyde. The product was triturated from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.46 (s, 1H), 7.56 (d, 2H, J 8.5 Hz) 7.45 (d, 2H, J 8.5 Hz), 7.11 (d, 1H, J 2.8 Hz), 7.04 (dd, 1H, J 8.9, 2.8 Hz), 6.96 (d, 1H, J 8.9 Hz), 5.58 (dd, 1H, J 13.1, 2.9 Hz), 3.15 (dd, 1H, J -16.8, 13.1 Hz), 2.79 (dd, 1H, J -16.8, 2.9 Hz).

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b) 2-(4-Chlorophenyl)chroman-4,6-diol

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2-(4-Chlorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 375 mg of 2-(4-chlorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.84 (s, 1H), 7.49-7.44 (m, 4H), 6.88 (d, 1H, J 2.8 Hz), 6.60 (d, 1H, J 8.6 Hz), 6.55 (dd, 1H, J 8.6, 2.8 Hz), 5.43 (bs, 1H), 5.14 (dd, 1H, J 11.9, 1.6 Hz), 4.87 (m, 1H), 2.26 (m, 1H), 1.85 (m, 1H).

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c) 2-(4-Chlorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

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2-(4-Chlorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 330 mg of 2-(4-chlorophenyl)chroman-4,6-diol. The product was purified by column chromatography using heptane- ethyl acetate (2:1) as an eluant. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.54-7.47 (m, 4H), 7.25 (d, 1H, J 2.8 Hz), 7.22 (d, 1H, 9.1 Hz), 7.02 (dd, 1H, J 8.8, 2.8 Hz), 6.89 (d, 1H, J 8.8 Hz), 5.65 (d, 1H, J 6.4 Hz), 5.33 (d, 1H, J 10.6 Hz), 4.98 (m, 1H), 2.34 (m, 1H), 1.94 (m, 1H).

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Example 69:

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2-[2-(2,4-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

a) 2-(2,4-Difluorophenyl)-6-hydroxychroman-4-one

2-(2,4-Difluorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 1,6 ml of 2,4-difluorobenzaldehyde. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.46 (s, 1H), 7.73 (m, 1H), 7.34 (m, 1H), 7.19 (m, 1H), 7.13 (d, 1H, J 2.9 Hz), 7.04 (dd, 1H, J 8.8, 2.9 Hz), 6.95 (d, 1H, J 8.8 Hz), 5.74 (dd, 1H, J 13.5, 2.8 Hz), 3.28 (dd, 1H, J -16.9, 13.5 Hz), 2.74 (dd, 1H, J -16.9, 2.8 Hz).

b) 2-(2,4-Difluorophenyl)chroman-4,6-diol

2-(2,4-Difluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1,47 g of 2-(2,4-difluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.86 (s, 1H), 7.61 (m, 1H), 7.28 (m, 1H), 7.14 (m, 1H), 6.88 (d, 1H, J 2.7 Hz), 6.59 (d, 1H, J 8.9 Hz), 6.54 (dd, 1H, J 8.9, 2.7 Hz), 5.46 (s, 1H), 5.32 (dd, 1H, J 11.9, 1.4 Hz), 4.88 (m, 1H), 2.24 (m, 1H), 1.99 (m, 1H).

c) 2-(2,4-Difluorophenyl)chroman-6-ol

2-(2,4-Difluorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 800 mg of 2-(2,4-difluorophenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.83 (s, 1H), 7.56 (m, 1H), 7.28 (m, 1H), 7.13 (m, 1H), 6.63 (m, 1H), 6.53-6.50 (m, 2H), 5.17 (dd, 1H, J 10.3, 2.3 Hz), 2.92 (ddd, 1H, J -17.0, 11.5, 5.8 Hz), 2.66 (ddd, 1H, J -17.0, 5.0, 2.7 Hz), 2.09 (m, 1H), 1.98 (m, 1H).

d) 2-[2-(2,4-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(2,4-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 720 mg of 2-(2,4-difluorophenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 3.0 Hz), 8.60 (dd, 1H, J 9.0, 3.0 Hz), 7.61 (m, 1H), 7.31 (m, 1H), 7.21 (d, 1H, 9.0 Hz), 7.17 (m, 1H), 7.02 (d, 1H, J 2.9 Hz), 6.97 (dd, 1H, J 8.9, 2.9 Hz), 6.91 (d, 1H, 8.9 Hz), 5.34 (dd, 1H, J 9.9, 2.0 Hz), 3.03 (m, 1H), 2.78 (m, 1H), 2.17 (m, 1H), 2.07 (m, 1H).

Example 70:**6-[2-(2,4-Difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine**

5 6-[2-(2,4-Difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 845 mg of 2-[2-(2,4-difluorophenyl)chroman-6-yloxy]-5-nitropyridine (Example 69(d)). ¹H NMR (400 MHz, d₆-DMSO) δ: 7.58 (m, 1H), 7.51 (d, 1H, J 3.3 Hz), 7.30 (m, 1H), 7.15 (m, 1H), 7.05 (dd, 1H, J 8.3, 3.3 Hz), 6.84-6.73 (m, 3H),
10 6.70 (d, 1H, J 8.3 Hz), 5.27 (dd, 1H, J 10.3, 2.3 Hz), 5.01 (s, 2H), 2.97 (m, 1H), 2.73 (m, 1H), 2.13 (m, 1H), 2.03 (m, 1H).

Example 71:

N-{6-[2-(2,4-Difluorophenyl)chroman-6-yloxy]pyridin-3-yl}methane sulfon-
15 **amide**

N-{6-[2-(2,4-difluorophenyl)chroman-6-yloxy]pyridin-3-yl} methane sulfonamide was prepared as described for N-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl] methane sulfonamide in Example 46 starting from 100 mg of 6-[2-(2,4-difluorophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 70). The product was crystallised
20 from a mixture of methanol and diethyl ether. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.67 (s, 1H), 7.99 (d, 1H, J 2.8 Hz), 7.67 (dd, 1H, J 8.8, 2.8 Hz), 7.60 (m, 1H), 7.30 (m, 1H), 7.16 (m, 1H), 6.98 (d, 1H, J 8.8 Hz) 6.92-6.86 (m, 3H), 5.31 (dd, 1H, J 10.3, 2.3 Hz), 3.01 (m, 1H), 2.98 (s, 3H), 2.76 (m, 1H), 2.16 (m, 1H), 2.06 (m, 1H).

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Example 72:**2-(2,4-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol**

2-(2,4-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared
30 as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 520 mg of 2-(2,4-difluorophenyl)chroman-4,6-diol (Example 69(b)). The product was recrystallised from a mixture of 2-propanol and diethyl ether. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.8 Hz), 8.61 (dd, 1H, J 9.1, 2.8 Hz), 7.66 (m, 1H), 7.32 (m, 1H), 7.26 (d, 1H, J 2.9 Hz), 7.23 (d, 1H, 9.1 Hz), 7.17 (m,
35 1H), 7.02 (dd, 1H, J 8.9, 2.9 Hz), 6.88 (d, 1H, J 8.9 Hz), 5.70 (bs, 1H), 5.52 (dd, 1H, J 11.9, 1.5 Hz), 5.00 (m, 1H), 2.31 (m, 1H), 2.09 (m, 1H).

Example 73**2-[2-(2-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine**

5 a) 2-(2-Chlorophenyl)-6-hydroxychroman-4-one

2-(2-Chlorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 2.8 g of 2-chlorobenzaldehyde. The product was
10 passed through silica gel using heptane - ethyl acetate as an eluant and then triturated with ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.49 (s, 1H), 7.77 (dd, 1H, J 7.7, 2.0 Hz), 7.53 (dd, 1H, J 7.6, 1.8 Hz), 7.49-7.41 (m, 2H), 7.14 (d, 1H, J 2.9 Hz), 7.06 (dd, 1H, J 8.8, 2.9 Hz), 6.93 (d, 1H, J 8.8 Hz), 5.78 (dd, 1H, J 13.6, 2.6 Hz), 3.19 (dd, 1H, J -16.9, 13.6 Hz), 2.78 (dd, 1H, J -16.9, 2.6 Hz).

15

b) 2-(2-Chlorophenyl)chroman-4,6-diol

2-(2-Chlorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.12 g of 2-(2-chlorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.63 (dd, 1H, J 7.7, 1.8 Hz), 7.49 (dd, 1H, J 7.8, 1.4 Hz), 7.45-7.36 (m, 2H), 6.89 (d, 1H, J 2.9 Hz), 6.63 (d, 1H, J 8.8 Hz), 6.56 (dd, 1H, J 8.9, 2.9 Hz), 5.39 (dd, 1H, J 11.7, 1.5 Hz), 4.90 (m, 1H), 2.33 (m, 1H), 1.82 (m, 1H).

25

c) 2-(2-Chlorophenyl)chroman-6-ol

2-(2-Chlorophenyl)chroman-6-ol I was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 500 mg of 2-(2-chlorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 7.58-7.36 (m, 4H), 6.66 (m, 1H), 6.55-6.51 (m, 2H), 5.23 (dd, 1H, J 10.1, 2.1 Hz), 2.92 (m, 1H), 2.68 (m, 1H),
30 2.17 (m, 1H), 1.87 (m, 1H).

d) 2-[2-(2-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine

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2-[2-(2-Chlorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting

from 485 mg of 2-(2-chlorophenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9, 0.5 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 7.62 (dd, 1H, J 7.5, 1.8 Hz), 7.51 (dd, 1H, J 7.6, 1.7 Hz), 7.45-7.40 (m, 2H), 7.21 (dd, 1H, J 9.1, 0.5 Hz), 7.04 (d, 1H, J 2.7 Hz), 6.99 (dd, 1H, J 8.8, 2.7 Hz), 6.94 (d, 1H, 8.8 Hz), 5.40 (dd, 1H, J 10.4, 2.1 Hz), 3.04 (m, 1H), 2.80 (m, 1H) 2.24 (m, 1H), 1.95 (m, 1H).

Example 74:**2-(2-Chlorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol**

2-(2-Chlorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 520 mg of 2-(2-chlorophenyl)chroman-4,6-diol (Example 68(b)). The product was recrystallised from 2-propanol. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.68 (dd, 1H, J 7.6, 1.8 Hz), 7.51-7.40 (m, 3H), 7.27 (d, 1H, J 2.9 Hz), 7.23 (d, 1H, J 9.1 Hz), 7.04 (dd, 1H, J 8.8, 2.9 Hz), 6.92 (d, 1H, J 8.8 Hz), 5.59 (d, 1H, J 10.6 Hz), 5.02 (m, 1H), 2.40 (m, 1H), 1.93 (m, 1H).

Example 75:**5-Nitro-2-[2-(4-Fluorophenyl)chroman-6-yloxy]pyridine**

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a) 6-Hydroxy-2-(4-fluorophenyl)chroman-4-one

6-Hydroxy-2-(4-fluorophenyl)chroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 2.0 g of 2',5'-dihydroxyacetophenone and 1.6 ml of 4-fluorobenzaldehyde. The product was recrystallised from acetic acid. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.59 (m, 2H), 7.27 (m, 2H), 7.14 (d, 1H, J 3.1 Hz), 7.05 (dd, 1H, J 8.9, 3.1 Hz), 6.96 (d, 1H, J 8.9 Hz), 5.56 (dd, 1H, J 13.2, 2.8 Hz), 3.18 (dd, 1H, J -16.9, 13.2 Hz), 2.77 (dd, 1H, J -16.9, 2.8 Hz).

30

b) 2-(4-Fluorophenyl)chroman-4,6-diol

2-(4-Fluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.5 g of 2-(4-fluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.84 (s, 1H), 7.48 (m, 2H),

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7.21 (m, 2H), 6.89 (d, 1H, J 2.7 Hz), 6.59 (d, 1H, J 8.7 Hz), 6.54 (dd, 1H, J 8.7, 2.7 Hz), 5.42 (bs, 1H), 5.12 (d, 1H, J 10.7 Hz), 4.87 (m, 1H), 2.25 (m, 1H), 1.89 (m, 1H).

c) 2-(4-Fluorophenyl)chroman-6-ol

5

2-(4-Fluorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 480 mg of 2-(4-fluorophenyl)-chroman-4,6-diol. ¹H NMR (400 MHz, CDCl₃) δ: 7.38 (m, 2H), 7.06 (m, 2H), 6.77 (d, 1H, J 8.6 Hz), 6.61 (dd, 1H, J 8.6, 2.9 Hz) 6.57 (d, 1H, 8.6 Hz), 4.97 (dd, 1H, J 10.2, 2.4 Hz), 2.95 (ddd, 1H, J -16.8, 11.4, 6.2 Hz), 2.74 (ddd, 1H, J -16.8, 5.3, 3.1 Hz), 2.15 (m, 1H), 2.05 (m, 1H).

10

d) 2-[2-(4-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine

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2-[2-(4-Fluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 160 mg of 2-(4-fluorophenyl)chroman-6-ol. The product was purified on preparative TLC-plate covered with silica gel using heptane- ethyl acetate (4:1) as an eluant. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (dd, 1H, J 2.9, 0.4 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 7.51 (m, 2H), 7.24 (m, 1H), 7.20 (dd, 1H, J 9.1, 0.4 Hz), 7.01 (d, 1H, J 2.8 Hz), 6.96 (dd, 1H, J 8.7, 2.8 Hz) 6.91 (d, 1H, 8.7 Hz), 5.15 (dd, 1H, J 10.3, 2.2 Hz), 2.94 (m, 1H), 2.76 (m, 1H) 2.17 (m, 1H), 2.01 (m, 1H).

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Example 76:

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6-[2-(4-Fluorophenyl)chroman-6-yloxy]pyridin-3-ylamine

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6-[2-(4-Fluorophenyl)chroman-6-yloxy]pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 3.04 g of 2-[2-(4-fluorophenyl)chroman-6-yloxy]-5-nitropyridine (Example 75(d)). ¹H NMR (400 MHz, d₆-DMSO) δ: 7.52-7.47 (m, 3H), 7.24 (m, 2H), 7.05 (dd, 1H, J 8.6, 3.0 Hz), 6.84-6.68 (m, 4H), 5.09 (dd, 1H, J 10.2, 2.1 Hz), 5.00 (bs, 2H), 2.93 (m, 1H), 2.69 (m, 1H), 2.13 (m, 1H), 1.98 (m, 1H).

Example 77:

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N-{6-[2-(4-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methane sulfonamide

N-{6-[2-(4-Fluorophenyl)chroman-6-yloxy]pyridin-3-yl}methane sulfonamide was prepared as described for N-[6-(2-phenylchroman-6-yloxy)pyridin-3-yl]methane sulfonamide in Example 46 starting from 442 mg of 6-[2-(4-fluorophenyl)-chroman-6-yloxy]-pyridin-3-ylamine (Example 76). The product was passed through silica gel using ethyl acetate - heptane (10:3) as an eluant and then crystallised from diethyl ether. ¹H NMR (400 MHz, CDCl₃) δ: 8.09 (d, 1H, J 2.8 Hz), 7.72 (dd, 1H, J 8.9, 2.8 Hz), 7.40 (m, 2H), 7.08 (m, 2H), 6.92-6.87 (m, 4H), 6.74 (s, 1H), 5.03 (dd, 1H, J 10.4, 2.3 Hz), 3.01 (m, 1H), 3.00 (s, 3H), 2.80 (m, 1H), 2.19 (m, 1H), 2.07 (m, 1H).

Example 78:**2-[2-(2,3-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine**

a) 2-(2,3-Difluorophenyl)-6-hydroxychroman-4-one

2-(2,3-Difluorophenyl)-6-hydroxychroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 2.6 ml of 2,3-difluorobenzaldehyde. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.51 (s, 1H), 7.53-7.46 (m, 2H), 7.31 (m, 1H), 7.14 (d, 1H, J 3.0 Hz), 7.05 (dd, 1H, J 8.8, 3.0 Hz), 6.96 (d, 1H, J 8.8 Hz), 5.82 (dd, 1H, J 13.4, 2.8 Hz), 3.26 (dd, 1H, J -16.9, 13.4 Hz), 2.79 (dd, 1H, J -16.9, 2.8 Hz).

b) 2-(2,3-Difluorophenyl)chroman-4,6-diol

2-(2,3-Difluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 2.91 g of 2-(2,3-difluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.88 (s, 1H), 7.45-7.36 (m, 2H), 7.28 (m, 1H), 6.89 (d, 1H, J 2.8 Hz), 6.61 (d, 1H, J 8.7 Hz), 6.55 (dd, 1H, J 8.7, 2.8 Hz), 5.49 (bs, 1H), 5.40 (dd, 1H, J 11.8, 1.4 Hz), 4.90 (m, 1H), 2.28 (m, 1H), 1.99 (m, 1H).

c) 2-(2,3-Difluorophenyl)chroman-6-ol

2-(2,3-Difluorophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 1.5 g of 2-(2,3-difluoro-

phenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 7.41 (m, 1H), 7.33 (m, 1H), 7.26 (m, 1H), 6.64 (dd, 1H, 9.0, 2.8 Hz), 6.54-6.51 (m, 2H), 5.25 (dd, 1H, J 10.2, 2.2 Hz), 2.93 (m, 1H), 2.66 (m, 1H), 2.14 (m, 1H), 2.01 (m, 1H).

5 d) 2-[2-(2,3-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(2,3-Difluorophenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 1.88 g of 2-(2,3-difluorophenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO)
10 δ: 9.04 (d, 1H, J 3.0 Hz), 8.60 (dd, 1H, J 9.1, 3.0 Hz), 7.45 (m, 1H), 7.38 (m, 1H), 7.30 (m, 1H), 7.21 (d, 1H, 9.1 Hz), 7.03 (d, 1H, J 2.7 Hz), 6.98 (dd, 1H, J 8.8, 2.7 Hz), 6.92 (d, 1H, 8.8 Hz), 5.42 (dd, 1H, J 10.4, 2.3 Hz), 3.04 (m, 1H), 2.79 (m, 1H) 2.21 (m, 1H), 2.08 (m, 1H).

15 **Example 79:**

2-(2,6-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

a) 6-Hydroxy-2-(2,6-difluorophenyl)chroman-4-one

20 6-Hydroxy-2-(2,6-Difluorophenyl)chroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 2.6 ml of 2,6-difluorobenzaldehyde. The product was triturated from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.55 (m, 1H) 7.22-7.18 (m, 2H), 7.14 (d, 1H, J 3.0 Hz), 7.03 (dd, 1H, J 8.9, 3.0 Hz), 6.93 (d, 1H, J
25 8.9 Hz), 5.84 (dd, 1H, J 14.0, 3.0 Hz), 3.38 (dd, 1H, J -17.0, 14.0 Hz), 2.80 (dd, 1H, J -17.0, 3.0 Hz).

b) 2-(2,6-Difluorophenyl)chroman-4,6-diol

30 2-(2,6-Difluorophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 4.45 g of 2-(2,6-difluorophenyl)-6-hydroxychroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.87 (s, 1H), 7.48 (m, 1H), 7.17-7.13 (m, 2H), 6.90 (d, 1H, J 2.9 Hz), 6.55-6.54 (m, 2H), 5.46 (dd, 1H, J 12.2, 1.8 Hz), 4.87 (m, 1H), 2.37 (m, 1H), 2.23 (m, 1H).

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c) 2-(2,6-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(2,6-Difluorophenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 1.9 g of 2-(2,6-difluorophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.03 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.52 (m, 1H), 7.26-7.16 (m, 4H), 7.01 (dd, 1H, J 8.8, 2.9 Hz), 6.83 (d, 1H, 8.8 Hz), 5.69-5.64 (m, 2H), 4.98 (m, 1H), 2.37 (m, 1H), 2.29 (m, 1H).

Example 80:

6-(5-Nitropyridin-2-yloxy)-2-(2-trifluoromethylphenyl)chroman-4-ol

a) 6-Hydroxy-2-(2-trifluoromethylphenyl)chroman-4-one

6-Hydroxy-2-(2-trifluoromethylphenyl)chroman-4-one was prepared as described for 2-(3-fluorophenyl)-6-hydroxychroman-4-one in Example 9(a) starting from 3.0 g of 2',5'-dihydroxyacetophenone and 3.0 ml of 2-trifluoromethylbenzaldehyde. The product was triturated from ethanol. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.48 (s, 1H), 8.07 (m, 1H), 7.86-7.79 (m, 2H), 7.66 (m, 1H), 7.15 (d, 1H, J 3.0 Hz), 7.06 (dd, 1H, J 8.8, 3.0 Hz), 6.95 (d, 1H, J 8.8 Hz), 5.70 (dd, 1H, J 13.8, 2.4 Hz), 3.38 (dd, 1H, J -16.9, 13.8 Hz), 2.66 (dd, 1H, J -16.9, 3.0 Hz).

b) 2-(2-Trifluoromethylphenyl)chroman-4,6-diol

2-(2-Trifluoromethylphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 1.43 g of 2-(2-trifluoromethylphenyl)-6-hydroxychroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.89 (s, 1H), 7.83 (m, 1H), 7.79-7.74 (m, 2H), 7.58 (m, 1H), 6.90 (d, 1H, J 2.7 Hz), 6.61 (d, 1H, J 8.9 Hz), 6.56 (dd, 1H, J 8.7, 2.7 Hz), 5.51 (d, 1H, J 6.5 Hz), 5.34 (d, 1H, J 11.6 Hz), 4.88 (m, 1H), 2.21 (m, 1H), 1.95 (m, 1H).

c) 6-(5-Nitropyridin-2-yloxy)-2-(2-trifluoromethylphenyl)chroman-4-ol

6-(5-Nitropyridin-2-yloxy)-2-(2-trifluoromethylphenyl)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 350 mg of 2-(2-trifluoromethylphenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.8 Hz), 8.62 (dd, 1H, J 9.1, 2.8 Hz), 7.89

(m, 1H), 7.82-7.78 (m, 2H), 7.62 (m, 1H), 7.28 (d, 1H, J 2.7 Hz), 7.24 (d, 1H, 9.1 Hz), 7.04 (dd, 1H, J 8.7, 2.7 Hz), 6.90 (d, 1H, J 8.7 Hz), 5.7 (bs, 1H), 5.38 (d, 1H, J 11.6 Hz), 5.01 (m, 1H), 2.29 (m, 1H), 2.05 (m, 1H).

5 **Example 81:**

2-[3-(3-Fluorophenyl)chroman-7-yloxy]-5-nitropyridine

a) 2-(3-Fluorophenyl)-1-(2-hydroxy-4-methoxyphenyl)ethanone

10 (3-Fluorophenyl)acetic acid (3.7 g) and 3-methoxyphenol (3.0 g) were dissolved into $\text{BF}_3 \cdot \text{Et}_2\text{O}$ (60 ml, 20 eq) under argon. The mixture was stirred at 60-70°C until disappearance of the starting materials (9 h) and poured into large volume of ice water. After extraction with ethyl acetate the combined organic layers were washed with water, dried and evaporated. The crude product was purified by column
15 chromatography using CH_2Cl_2 as an eluant. ^1H NMR (400 MHz, d_6 -DMSO) δ : 12.41 (br s, 1H), 8.02 (d, 1H, J 9.0 Hz), 7.34-7.38 (m, 1H), 7.09-7.13 (m, 3H), 6.56 (dd, 1H, J 9.0, 2.5 Hz), 6.49 (d, 1H, J 2.5 Hz), 4.41 (s, 2H), 3.83 (s, 3H).

b) 3-(3-Fluorophenyl)-7-methoxychromen-4-one

20 2-(3-Fluorophenyl)-1-(2-hydroxy-4-methoxyphenyl)ethanone (1.76 g) was dissolved in pyridine (88 ml). Piperidine (8.8 ml) and triethylorthoformate (88 ml) were added and the mixture was stirred at 120°C for 3.5 hours. After pouring the mixture into water and acidification with conc. HCl the crude product was filtered.
25 Purification by column chromatography using heptane-ethyl acetate (7:3) as an eluant afforded 3-(3-fluorophenyl)-7-methoxychromen-4-one. ^1H NMR (400 MHz, d_6 -DMSO) δ : 8.57 (s, 1H), 8.06 (d, 1H, J 8.9 Hz), 7.45-7.50 (m, 3H), 7.21-7.25 (m, 1H), 7.20 (d, 1H, J 2.4 Hz), 7.12 (dd, 1H, J 8.9, 2.4 Hz), 3.92 (s, 3H).

30 c) 3-(3-Fluorophenyl)-7-hydroxychromen-4-one

3-(3-Fluorophenyl)-7-methoxychromen-4-one (320 mg) was refluxed with 47 % HBr (18 ml) until disappearance of the starting material. The mixture was poured into water and the precipitate was filtrated and dried yielding 3-(3-fluorophenyl)-7-
35 hydroxychromen-4-one. ^1H NMR (400 MHz, d_6 -DMSO) δ : 10.87 (s, 1H), 8.49 (s,

1H), 7.99 (d, 1H, J 8.7 Hz), 7.43-7.49 (m, 3H), 7.20-7.24 (m, 1H), 6.97 (dd, 1H, J 8.7, 2.2 Hz), 6.90 (d, 1H, J 2.2 Hz).

d) 3-(3-Fluorophenyl)chroman-7-ol

5

3-(3-Fluorophenyl)-7-hydroxychromen-4-one (160 mg) was dissolved in ethanol (40 ml) and 10 % palladium on carbon (400 mg) was added. The reaction mixture was hydrogenated for 6 hours at normal pressure and room temperature. It was then filtered through Celite and washed with ethanol. The solvent was evaporated under reduced pressure to give 3-(3-fluorophenyl)chroman-7-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.19 (br s, 1H), 7.38 (m, 1H), 7.17-7.21 (m, 2H), 7.08 (m, 1H), 6.88 (d, 1H, J 8.2 Hz), 6.30 (dd, 1H, J 8.2, 2.4 Hz), 6.20 (d, 1H, J 2.4 Hz), 4.22 (dd, 1H, J 10.3, 3.6 Hz), 4.02 (t, 1H, 10.3 Hz), 3.20 (m, 1H), 2.90 (m, 2H).

10

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e) 2-[3-(3-Fluorophenyl)chroman-7-yloxy]-5-nitropyridine

2-[3-(3-Fluorophenyl)chroman-7-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 125 mg of 3-(3-fluorophenyl)-chroman-7-ol. The product was recrystallised from ethanol. ¹H NMR (400 MHz, CDCl₃) δ: 9.07 (d, 1H, J 2.8 Hz), 8.47 (dd, 1H, J 9.0, 2.8 Hz), 7.33 (m, 1H), 7.16 (d, 1H J 8.9 Hz), 6.95-7.06 (m, 4H), 6.69-6.71 (m, 2H), 4.38 (dd, 1H, J 10.6, 4.3 Hz), 4.06 (t, 1H, 10.6 Hz), 3.30 (m, 1H), 3.06 (m, 2H).

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Example 82:

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5-Nitro-2-(3-phenylchroman-7-yloxy)pyridine

a) 7-Hydroxy-3-phenylchromen-4-one

7-Hydroxy-3-phenylchromen-4-one is commercially available or can be synthesised by methods described for 3-(3-fluorophenyl)-7-hydroxychromen-4-one (Example 81(a-c)). ¹H NMR spectrum as reported in the literature (*Synth. Commun.*, 2000, 30(3), 469-484).

30

b) 3-Phenylchroman-7-ol

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3-Phenylchroman-7-ol was prepared as described for 3-(3-fluorophenyl)-chroman-7-ol in Example 81(d) using 0.5 g of 7-hydroxy-3-phenylchromen-4-one.
¹H NMR (400 MHz, d₆-DMSO) δ: 8.18 (br s, 1H), 7.31-7.34 (m, 4H), 7.25-7.27 (m, 1H), 6.88 (d, 1H, J 8.2 Hz), 6.30 (dd, 1H, J 8.2, 2.4 Hz), 6.20 (d, 1H, J 2.4 Hz), 4.21
5 (dd, 1H, J 10.3, 3.6 Hz), 4.00 (t, 1H, 10.3 Hz), 3.13 (m, 1H), 2.84-2.87 (m, 2H).

c) 5-Nitro-2-(3-phenylchroman-7-yloxy)pyridine

5-Nitro-2-(3-phenylchroman-7-yloxy)pyridine was prepared as described for
10 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) using 200 mg of 3-phenylchroman-7-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.05 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.34-7.38 (m, 4H), 7.27-7.30 (m, 1H), 7.22 (m, 2H), 6.70-6.74 (m, 2H), 4.31 (dd, 1H, J 10.4, 3.5 Hz), 4.12 (t, 1H, 10.4 Hz), 3.24 (m, 1H), 3.01-3.11 (m, 2H).

15

Example 83:

5-Methoxy-2-(3-phenylchroman-7-yloxy)phenylamine hydrochloride

a) 7-(4-Methoxy-2-nitrophenoxy)-3-phenylchroman

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7-(4-Methoxy-2-nitrophenoxy)-3-phenylchroman was prepared as described for 6-(4-methoxy-2-nitrophenoxy)-2-phenylchroman in Example 3(a) using 0.5 g of 3-phenylchroman-7-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.60 (d, 1H, J 3.1 Hz), 7.35 (m, 4H), 7.32 (dd, 1H, J 9.1, 3.1 Hz), 7.27-7.29 (m, 1H), 7.21 (d, 1H, J 9.1 Hz),
25 7.12 (d, 1H, J 8.3 Hz), 6.48 (dd, 1H, J 8.3, 2.5 Hz), 6.38 (d, 1H, J 2.5 Hz), 4.26 (dd, 1H, J 10.4, 3.5 Hz), 4.08 (t, 1H, J 10.4 Hz), 3.85 (s, 3H), 3.20 (m, 1H), 2.90-3.04 (m, 2H).

b) 5-Methoxy-2-(3-phenylchroman-7-yloxy)phenylamine hydrochloride

30

5-Methoxy-2-(3-phenylchroman-7-yloxy)phenylamine hydrochloride was prepared as described for 5-methoxy-2-(2-phenylchroman-6-yloxy)phenylamine hydrochloride in Example 3(b) using 310 mg of 7-(4-methoxy-2-nitrophenoxy)-3-phenylchroman. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.32-7.35 (m, 4H), 7.23-7.29 (m,
35 1H), 7.05 (d, 1H, J 8.4 Hz), 6.76 (d, 1H, J 8.8 Hz), 6.46 (d, 1H, J 2.8 Hz), 6.43 (dd,

1H, J 8.4, 2.5 Hz), 6.28 (d, 1H, J 2.5 Hz), 6.24 (dd, 1H, J 8.8, 2.8 Hz), 4.24 (dd, 1H, J 10.4, 3.4 Hz), 4.05 (t, 1H, J 10.4 Hz), 3.18 (m, 1H), 2.88-3.01 (m, 2H).

Example 84:

5 5-Nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine

a) 2-(2-Hydroxy-1-phenylethylsulfanyl)benzene-1,4-diol

To a stirred solution of 2-mercaptobenzene-1,4-diol (0.5 g) and potassium
10 carbonate (0.49 g) in water (5 ml) was added 2-phenyloxirane (0.40 ml) under argon.
The mixture was stirred at room temperature for 2.5 hours and then treated with 2 M
HCl and extracted with ethyl acetate. The combined organic layers were washed
with water and brine, dried and evaporated. The crude product was purified by
column chromatography using heptane-ethyl acetate (1:1) as an eluant. ¹H NMR (400
15 MHz, d₆-DMSO) δ: 8.94 (br s, 1H), 8.72 (br s, 1H), 7.24-7.37 (m, 5H), 6.62-6.65 (m,
2H), 6.47 (dd, 1H, J 8.6, 2.8 Hz), 4.97 (br s, 1H), 4.34 (m, 1H), 3.72 (m, 2H).

b) 2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-ol

20 A solution of 2-(2-hydroxy-1-phenylethylsulfanyl)benzene-1,4-diol (0.83 g) in
dry toluene (60 ml) was stirred with Amberlyst 15 (0.5 g) at 60 °C until disappear-
ance of the starting material. After the mixture was filtered and solvent evaporated
the crude product was purified by column chromatography using heptane-ethyl
acetate (1:1) as an eluant. ¹H NMR (400 MHz, CDCl₃) δ: 7.41 (m, 4H), 7.33-7.40
25 (m, 1H), 6.81 (d, 1H, J 8.7 Hz), 6.61 (d, 1H, J 3.0 Hz), 6.51 (dd, 1H, J 8.7, 3.0 Hz),
5.10 (dd, 1H, J 9.6, 1.9 Hz), 3.28 (dd, 1H, J 13.0, 9.6 Hz), 3.06 (dd, 1H, J 13.0, 1.9
Hz).

c) 5-Nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine

30 5-Nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine was
prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example
1(b) using 269 mg 2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-ol. The product was
recrystallised from ethanol. ¹H NMR (400 MHz, CDCl₃) δ: 9.07 (d, 1H, J 2.8 Hz),
35 8.47 (dd, 1H, J 9.1, 2.8 Hz), 7.43 (m, 4H), 7.37-7.41 (m, 1H), 7.02 (d, 1H, J 9.1 Hz),

6.99 (d, 1H, J 8.9 Hz), 6.95 (d, 1H, J 2.8 Hz), 6.82 (dd, 1H, J 8.9, 2.8 Hz), 5.21 (dd, 1H, J 9.7, 1.9 Hz), 3.31 (dd, 1H, 13.2, 9.7 Hz), 3.11 (dd, 1H, 13.2, 1.9 Hz).

Example 85:

5 5-Nitro-2-(4-oxo-2-phenyl-3,4-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine

To 5-nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine (214 mg) in methanol (80 ml) at 60°C, NaIO₄ (total 2.5 eq) was added in small portions until disappearance of the starting material. The mixture was poured into water and precipitate was filtered and washed with water. The crude product was purified by column chromatography using heptane-ethyl acetate (1:1) as an eluant. ¹H NMR (400 MHz, CDCl₃) δ: 9.02 (d, 1H, J 2.8 Hz), 8.51 (dd, 1H, J 9.1, 2.8 Hz), 7.45-7.56 (m, 6H), 7.32 (dd, 1H, J 9.0, 2.8 Hz), 7.21 (d, 1H, J 9.0 Hz), 7.10 (d, 1H, J 9.1 Hz), 5.77 (dd, 1H, J 12.0, 1.5 Hz), 3.34 (dd, 1H, 14.4, 1.5 Hz), 3.13 (dd, 1H, 14.4, 12.0 Hz).

15

Example 86:

2-(4,4-Dioxo-2-phenyl-3,4-dihydro-2-benzo[1,4]oxathiin-6-yloxy)-5-nitropyridine

20 To stirred solution of 5-nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine (150 mg) in AcOH (3.7 ml) and water (1.2 ml) was added KMnO₄ (125 mg) in an ice/water bath. The mixture was stirred at room temperature for 1.5 hours, then water (5ml) and 30% H₂O₂ were added until decomposition of excess KMnO₄ was complete. The mixture was diluted with additional water (20 ml). The white precipitate was filtered, washed with water and dried to give 2-(4,4-dioxo-2-phenyl-3,4-dihydro-2-benzo[1,4]oxathiin-6-yloxy)-5-nitropyridine. ¹H NMR (400 MHz, CDCl₃) δ: 9.03 (d, 1H, J 2.8 Hz), 8.52 (dd, 1H, J 9.0, 2.8 Hz), 7.68 (d, 1H, J 2.8 Hz), 7.46-7.52 (m, 5H), 7.33 (dd, 1H, J 9.1, 2.8 Hz), 7.17 (d, 1H, J 9.1 Hz), 7.11 (d, 1H, J 9.0 Hz), 5.87 (dd, 1H, J 12.2, 1.6 Hz), 3.76 (dd, 1H, 14.1, 12.2 Hz), 3.55 (dd, 1H, 14.1, 1.6 Hz).

30

Example 87:

5-Nitro-2-[2-(4-nitrophenyl)chroman-6-yloxy]pyridine

35 a) 6-Hydroxy-2-(4-nitrophenyl)chroman-4-one

6-Hydroxy-2-(4-nitrophenyl)chroman-4-one was prepared as described for 6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was recrystallised from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.48 (s, 1H), 8.29 (d, 2H, J 6.9 Hz), 7.83 (d, 2H, J 6.9 Hz), 7.13 (d, 1H J 2.9 Hz), 7.06 (dd, 1H, J 8.8, 2.9 Hz), 7.01 (d, 1H, J 8.8 Hz), 5.77 (dd, 1H, J 13.0, 3.0 Hz), 3.15 (dd, 1H, J 16.8, 13.0 Hz), 2.89 (dd, 1H, J 16.8, 3.0 Hz).

b) 2-(4-Nitrophenyl)chroman-4,6-diol

2-(4-Nitrophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(4-nitrophenyl)chroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.86 (s, 1H), 8.26 (d, 2H, J 6.9 Hz), 7.74 (d, 2H, J 6.9 Hz), 6.89 (d, 1H J 2.8 Hz), 6.65 (d, 1H, J 8.6 Hz), 6.56 (dd, 1H, J 8.6, 2.8 Hz), 5.46 (d, 1H, J 6.9 Hz), 5.32 (d, 1H, J 10.5 Hz), 4.86-4.94 (m, 1H), 2.31-2.38 (m, 1H), 1.78-1.89 (m, 1H).

c) 2-(4-Nitrophenyl)chroman-6-ol

2-(4-Nitrophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 2-(4-nitrophenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.84 (s, 1H), 8.26 (d, 2H, J 6.9 Hz), 7.71 (d, 2H, J 6.9 Hz), 6.69 (d, 1H, J 8.6 Hz), 6.53 (dd, 1H, J 8.6, 2.8 Hz), 6.50 (d, 1H, J 2.8 Hz), 5.19 (dd, 1H, J 9.9, 2.2 Hz), 2.87-2.91 (m, 1H), 2.61-2.66 (m, 1H), 2.16-2.21 (m, 1H), 1.89-1.93 (m, 1H).

d) 5-Nitro-2-[2-(4-nitrophenyl)chroman-6-yloxy]pyridine

5-Nitro-2-[2-(4-nitrophenyl)chroman-6-yloxy]pyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(4-nitrophenyl)chroman-6-ol. ¹H NMR (300 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 8.29 (d, 2H, J 6.9 Hz), 7.76 (d, 2H, J 6.9 Hz), 7.21 (d, 1H, J 9.1 Hz), 6.98-7.02 (m, 3H), 5.35 (dd, 1H, J 9.9, 2.2 Hz), 2.96-3.05 (m, 1H), 2.73-2.78 (m, 1H), 2.24-2.29 (m, 1H), 1.96-2.04 (m, 1H).

Example 88:**6-[2-(4-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine**

6-[2-(4-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as
5 described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using
100 mg 5-nitro-2-[2-(4-nitrophenyl)chroman-6-yloxy]pyridine (Example 87(d)) and
665 mg of Zn. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.50 (d, 1H, J 2.9 Hz), 7.07 (d, 2H,
8.4 Hz), 7.04 (dd, 1H, J 8.6, 2.9 Hz), 6.71 (s, 3H), 6.68 (d, 1H, J 8.6 Hz), 6.56 (d, 2H,
J 8.4 Hz), 5.07 (s, 2H), 4.99 (s, 2H), 4.84 (dd, 1H, J 9.7, 2.3 Hz), 2.86-2.95 (m, 1H),
10 2.66-2.71 (m, 1H), 1.95-2.05 (m, 2H).

Example 89:**5-Nitro-2-[2-(2-nitrophenyl)chroman-6-yloxy]pyridine**

15 a) 6-Hydroxy-2-(2-nitrophenyl)chroman-4-one

6-Hydroxy-2-(2-nitrophenyl)chroman-4-one was prepared as described for 6-
hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). ¹H NMR (400 MHz, d₆-
DMSO) δ: 9.49 (s, 1H), 8.05-8.06 (m, 1H), 7.96-7.98 (m, 1H), 7.83-7.87 (m, 1H),
20 7.65-7.69 (m, 1H), 7.14 (d, 1H, J 3.1 Hz), 7.05 (dd, 1H, J 8.8, 3.1 Hz), 6.91 (d, 1H, J
8.8 Hz), 5.69 (dd, 1H, J 13.0, 2.6 Hz), 3.22 (dd, 1H, J 16.8, 13.0 Hz), 2.98 (dd, 1H, J
16.8, 2.6 Hz).

b) 2-(2-Nitrophenyl)chroman-4,6-diol

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2-(2-Nitrophenyl)chroman-4,6-diol was prepared as described for 2-phenyl-
chroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(2-nitrophenyl)-
chroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.87 (s, 1H), 7.99-8.02 (m, 1H),
7.77-7.86 (m, 2H), 7.59-7.64 (m, 1H), 6.89 (d, 1H, J 2.4 Hz), 6.56-6.57 (m, 2H),
30 5.51-5.55 (m, 2H), 4.85-4.92 (m, 1H), 2.42-2.47 (m, 1H), 1.85-1.96 (m, 1H).

c) 2-(2-Nitrophenyl)chroman-6-ol

2-(2-Nitrophenyl)chroman-6-ol was prepared as described for 2-(3-fluoro-
35 phenyl)chroman-6-ol in Example 9(c) starting from 2-(2-nitrophenyl)chroman-4,6-
diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.85 (s, 1H), 8.00 (d, 1H, J 8.0 Hz), 7.79-

7.80 (m, 2H), 7.59-7.63 (m, 1H), 6.59-6.62 (m, 1H), 6.50-6.53 (m, 2H), 5.36 (dd, 1H, J 10.2, 2.0 Hz), 2.89-2.93 (m, 1H), 2.67-2.73 (m, 1H), 2.26-2.31 (m, 1H), 1.90-1.95 (m, 1H).

5 d) 5-Nitro-2-[2-(2-nitrophenyl)chroman-6-yloxy]pyridine

5-Nitro-2-[2-(2-nitrophenyl)chroman-6-yloxy]pyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(2-nitrophenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 8.03 (d, 1H, J 7.9 Hz), 7.80-7.85 (m, 2H), 7.62-7.66 (m, 1H), 7.22 (d, 1H, J 9.1 Hz), 7.04 (d, 1H, J 2.8 Hz), 6.98 (dd, 1H, J 8.8, 2.8 Hz), 6.88 (d, 1H, J 8.8 Hz), 5.52 (dd, 1H, J 10.3, 2.0 Hz), 2.99-3.31 (m, 1H), 2.80-2.85 (m, 1H), 2.35-2.40 (m, 1H), 1.99-2.04 (m, 1H).

15 **Example 90:**

6-[2-(2-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine

6-[2-(2-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using 20 100 mg 5-nitro-2-[2-(2-nitrophenyl)chroman-6-yloxy]pyridine (Example 89(d)) and 700 mg of Zn. ¹H NMR (300 MHz, d₆-DMSO) δ: 7.51 (d, 1H, J 2.9 Hz), 7.15-7.18 (m, 1H), 7.05 (dd, 1H, J 8.6, 2.9 Hz), 6.98-7.00 (m, 1H), 6.77 (d, 1H, J 8.6 Hz), 6.73-6.75 (m, 2H), 6.66-6.71 (m, 2H), 6.56-6.61 (m, 1H), 5.11 (dd, 1H, J 10.4, 2.0 Hz), 5.01 (s, 2H), 4.99 (s, 2H), 2.94-2.99 (m, 1H), 2.66-2.74 (m, 1H), 2.06-2.13 (m, 1H), 25 1.88-1.95 (m, 1H).

Example 91:

N-{6-[2-(2-Acetylaminophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

30 N-{6-[2-(2-Acetylaminophenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetyl-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 6-[2-(2-aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine (Example 90). ¹H NMR (300 MHz, d₆-DMSO) δ: 10.02 (s, 1H), 9.44 (s, 1H), 8.27 (d, 1H, J 2.7 Hz), 8.00 (dd, 1H, J 8.8, 2.7 Hz), 7.49 (d, 1H, J 8.8 Hz), 7.25-7.38 (m, 3H), 6.92 (d, 1H, J 8.9 Hz), 6.88 (s, 1H), 6.83 (s, 2H), 5.22 (d, 1H, J 8.7 Hz), 2.90-2.99 35 (m, 1H), 2.72-2.79 (m, 1H), 2.12-2.18 (m, 1H), 2.04 (s, 6H), 1.86-1.94 (m, 1H).

Example 92:**5-Nitro-2-[2-(3-nitrophenyl)chroman-6-yloxy]pyridine**

5 a) 6-Hydroxy-2-(3-nitrophenyl)chroman-4-one

6-Hydroxy-2-(3-nitrophenyl)chroman-4-one was prepared as described for 6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was recrystallised from ethanol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.40 (s, 1H), 8.24 (dd, 1H, J 8.2, 2.3 Hz), 8.01 (d, 1H, J 7.9 Hz), 7.74 (t, 1H, J 15.9, 7.9 Hz), 7.13 (d, 1H, J 2.9 Hz), 7.07 (dd, 1H, J 8.8, 2.9 Hz), 7.00 (d, 1H, 8.8 Hz), 5.75 (dd, 1H, J 13.1, 2.9 Hz), 3.21 (dd, 1H, J 16.8, 13.1 Hz), 2.88 (dd, 1H, J 16.8, 2.9 Hz).

15 b) 2-(3-Nitrophenyl)chroman-4,6-diol

2-(3-Nitrophenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(3-nitrophenyl)chroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.89 (br s, 1H), 8.29 (s, 1H), 8.20 (dd, 1H, J 8.2, 2.3 Hz), 7.93 (d, 1H, J 7.9 Hz), 7.71 (t, 1H, J 15.9, 7.9 Hz), 6.89 (d, 1H, J 2.8 Hz), 6.66 (d, 1H, J 8.7 Hz), 6.57 (dd, 1H, J 8.7, 2.9 Hz), 5.47 (br s, 1H), 5.33 (d, 1H, J 10.7 Hz), 4.88-4.92 (m, 1H), 2.33-2.39 (m, 1H), 1.83-1.92 (m, 1H).

c) 2-(3-Nitrophenyl)chroman-6-ol

25 2-(3-Nitrophenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 2-(3-nitrophenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.80 (s, 1H), 8.26 (s, 1H), 8.19 (dd, 1H, J 8.1, 2.3 Hz), 7.90 (d, 1H, J 7.9 Hz), 7.70 (t, 1H, J 15.9, 7.9 Hz), 6.70 (d, 1H, J 8.4 Hz), 6.51-6.55 (m, 2H), 5.19 (dd, 1H, J 10.0, 2.0), 2.86-2.91 (m, 1H), 2.61-2.68 (m, 1H), 2.17-2.23 (m, 1H), 1.91-1.97 (m, 1H).

d) 5-Nitro-2-[2-(3-nitrophenyl)chroman-6-yloxy]pyridine

35 5-Nitro-2-[2-(3-nitrophenyl)chroman-6-yloxy]pyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(3-nitrophenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H,

J 2.9 Hz), 8.60 (dd, 1H, J 9.0, 2.9 Hz), 8.32 (s, 1H), 8.23 (d, 1H, J 8.3 Hz), 7.95 (d, 1H, J 7.9 Hz), 7.74 (t, 1H, J 15.8, 7.9 Hz), 7.21 (d, 1H, J 9.0 Hz), 6.96-7.03 (m, 3H), 5.35 (d, 1H, J 8.7 Hz), 2.98-3.06 (m, 1H), 2.72-2.79 (m, 1H), 2.26-2.33 (m, 1H), 1.99-2.06 (m, 1H).

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Example 93:**6-[2-(3-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine**

6-[2-(3-Aminophenyl)chroman-6-yloxy]-pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using 150 mg 5-nitro-2-[2-(3-nitrophenyl)chroman-6-yloxy]pyridine (Example 92(d)) and 997 mg of Zn. ¹H NMR (300 MHz, d₆-DMSO) δ: 7.51 (d, 1H, J 2.8 Hz), 7.05 (dd, 1H, J 8.6, 2.8 Hz), 7.01 (t, 1H, J 15.4, 7.7 Hz), 6.70-6.78 (m, 3H), 6.68 (d, 1H, J 8.6 Hz), 6.63 (s, 1H), 6.54 (d, 1H, J 7.7 Hz), 6.50 (d, 1H, J 8.6 Hz), 5.06 (s, 2H), 4.98 (s, 2H), 4.90 (dd, 1H, J 10.0, 2.2 Hz), 2.85-2.96 (m, 1H), 2.62-2.74 (m, 1H), 2.05-2.11 (m, 1H), 1.89-1.95 (m, 1H).

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Example 94:**2-(4-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol**

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a) 6-Hydroxy-2-(4-methoxyphenyl)chroman-4-one

6-Hydroxy-2-(4-methoxyphenyl)chroman-4-one was prepared as described for 6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was recrystallised from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.40 (s, 1H), 7.45 (d, 2H, J 8.7 Hz), 7.11 (d, 1H, J 3.0 Hz), 7.02 (dd, 1H, J 8.9, 3.0 Hz), 6.97 (d, 2H, J 8.7 Hz), 6.93 (d, 1H, J 8.9 Hz), 5.47 (dd, 1H, J 13.1, 2.8 Hz), 3.77 (s, 3H), 3.19 (dd, 1H, J 16.8, 13.1 Hz), 2.72 (dd, 1H, J 16.8, 2.8 Hz).

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b) 2-(4-methoxyphenyl)chroman-4,6-diol

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2-(4-Methoxyphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(4-methoxyphenyl)chroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.78 (s, 1H), 7.35 (d, 2H, J 8.7 Hz), 6.94 (d, 2H, J 8.7 Hz), 6.88 (d, 1H, J 2.5 Hz), 6.56 (d, 1H, J 8.7 Hz), 6.52

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(dd, 1H, J 8.7, 2.5 Hz), 5.37 (br s, 1H), 5.04 (d, 1H, J 10.9 Hz), 4.83-4.89 (m, 1H), 3.76 (s, 3H), 2.18-2.25 (m, 1H), 1.85-1.97 (m, 1H).

c) 2-(4-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

5

2-(4-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(4-methoxyphenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.40 (d, 2H, J 8.7 Hz), 7.24 (d, 1H, J 2.8 Hz), 7.22 (d, 1H, J 9.1 Hz), 7.00 (dd, 1H, J 8.7, 2.8 Hz), 6.97 (d, 2H, J 8.7 Hz), 6.84 (d, 1H, J 8.7 Hz), 5.63 (d, 1H, J 6.4 Hz), 5.23 (d, 1H, J 10.8 Hz), 4.95-5.02 (m, 1H), 3.78 (s, 3H), 2.25-2.29 (m, 1H), 1.98-2.04 (m, 1H).

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Example 95:

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6-(5-Aminopyridin-2-yloxy)-2-(4-methoxyphenyl)chroman-4-ol

6-(5-Aminopyridin-2-yloxy)-2-(4-methoxyphenyl)chroman-4-ol was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using 105 mg 2-(4-methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol (Example 94(c)) and 348 mg of Zn. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.52 (d, 1H, J 3.0 Hz), 7.38 (d, 2H, J 8.8 Hz), 7.06 (dd, 1H, J 8.7, 3.0 Hz), 7.04 (d, 1H, J 2.9 Hz), 6.96 (d, 2H, J 8.8 Hz), 6.77 (dd, 1H, J 8.7, 2.9 Hz), 6.72 (d, 1H, J 8.7 Hz), 6.70 (d, 1H, J 8.6 Hz), 5.52 (d, 1H, J 6.6 Hz), 5.15 (d, 1H, J 10.7 Hz), 5.01 (s, 2H), 4.87-4.93 (m, 1H), 3.77 (s, 3H), 2.22-2.26 (m, 1H), 1.90-1.99 (m, 1H).

25

Example 96:

N-{6-[4-Hydroxy-2-(4-methoxyphenyl)chroman-6-yloxy]pyridin-3-yl}
acetamide

30

N-{6-[4-Hydroxy-2-(4-methoxyphenyl)chroman-6-yloxy]pyridin-3-yl}-acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 6-(5-aminopyridin-2-yloxy)-2-(4-methoxyphenyl)chroman-4-ol (Example 95). ¹H NMR (400 MHz, d₆-DMSO) δ: 10.05 (s, 1H), 8.28 (d, 1H, J 2.7 Hz), 8.01 (dd, 1H, J 8.8, 2.7 Hz), 7.39 (d, 2H, J 8.7 Hz), 7.14 (d, 1H, J 2.7 Hz), 6.97 (d, 2H, J 8.7 Hz), 6.94 (d, 1H, J 8.8 Hz), 6.88 (dd, 1H, J 8.8,

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2.7 Hz), 6.77 (d, 1H, J 8.8 Hz), 5.57 (d, 1H, J 6.5 Hz), 5.19 (d, 1H, J 10.6 Hz), 4.90-4.97 (m, 1H), 3.77 (s, 3H), 2.24-2.27 (m, 1H), 2.04 (s, 3H), 1.93-2.01 (m, 1H).

Example 97:

5 2-(2-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

a) 6-Hydroxy-2-(2-methoxyphenyl)chroman-4-one

6-Hydroxy-2-(2-methoxyphenyl)chroman-4-one was prepared as described for
10 6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was
recrystallised from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.40 (s, 1H), 7.56 (dd,
1H, J 7.6, 1.6 Hz), 7.37 (dt, 1H, J 8.6, 7.0, 1.6 Hz), 7.12 (d, 1H, J 2.9 Hz), 7.07 (d,
1H, J 8.6 Hz), 7.03 (d, 1H, J 8.9 Hz), 7.02 (d, 1H, J 7.0 Hz), 6.95 (d, 1H, J 8.9 Hz),
5.70 (dd, 1H, J 13.3, 2.7 Hz), 3.82 (s, 3H), 3.09 (dd, 1H, J 16.8, 13.3 Hz), 2.71 (dd,
15 1H, J 16.8, 2.7 Hz).

b) 2-(2-methoxyphenyl)chroman-4,6-diol

2-(2-Methoxyphenyl)chroman-4,6-diol was prepared as described for 2-phe-
20 nylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(2-methoxyphenyl)-
chroman-4-one. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.79 (s, 1H), 7.45 (dd, 1H, J 7.6,
1.6 Hz), 7.31 (dt, 1H, J 8.5, 7.3, 1.6 Hz), 7.04 (d, 1H, J 8.5 Hz), 6.99 (d, 1H, J 7.3
Hz), 6.88 (d, 1H, J 2.7 Hz), 6.59 (d, 1H, J 8.7 Hz), 6.54 (dd, 1H, J 8.7, 2.7 Hz), 5.38
(s, 1H), 5.34 (d, 1H, J 11.4 Hz), 4.80-4.88 (m, 1H), 3.81 (s, 3H), 2.24-2.28 (m, 1H),
25 1.74-1.86 (m, 1H).

c) 2-(2-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol

2-(2-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared
30 as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b)
starting from 2-(2-methoxyphenyl)chroman-4,6-diol. ¹H NMR (400 MHz, d₆-
DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, J 9.1, 2.9 Hz), 7.49 (dd, 1H, J 7.6,
1.7 Hz), 7.34 (dt, 1H, J 8.3, 7.5, 1.7 Hz), 7.24 (d, 1H, J 2.8 Hz), 7.22 (d, 1H, J 9.1
Hz), 7.07 (d, 1H, J 8.3 Hz), 7.03 (d, 1H, J 7.5 Hz), 7.01 (dd, 1H, J 8.7, 2.8 Hz), 6.88
35 (d, 1H, J 8.7 Hz), 5.62 (d, 1H, J 6.3 Hz), 5.52 (d, 1H, J 10.4 Hz), 4.93-4.99 (m, 1H),
3.84 (s, 3H), 2.30-2.35 (m, 1H), 1.85-1.94 (m, 1H).

Example 98:**6-(5-Aminopyridin-2-yloxy)-2-(2-methoxyphenyl)chroman-4-ol**

6-(5-Aminopyridin-2-yloxy)-2-(2-methoxyphenyl)chroman-4-ol was prepared
5 as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using
79 mg 2-(2-methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol (Example
97(c)) and 262 mg of Zn. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.52 (d, 1H, J 2.9 Hz),
7.47 (dd, 1H, J 7.5, 1.6 Hz), 7.33 (dt, 1H, J 8.5, 7.4, 1.6 Hz), 7.06 (d, 1H, J 2.8 Hz),
7.03-7.05 (m, 2H), 7.01 (d, 1H, J 7.4 Hz), 6.79 (dd, 1H, J 8.6, 2.8 Hz), 6.75 (d, 1H, J
10 8.7 Hz), 6.71 (d, 1H, J 8.6 Hz), 5.52 (d, 1H, J 6.5 Hz), 5.44 (d, 1H, J 10.5 Hz), 5.01
(s, 2H), 4.86-4.92 (m, 1H), 3.83 (s, 3H), 2.27-2.32 (m, 1H), 1.80-1.88 (m, 1H).

Example 99:**2-[2-(3-Methoxyphenyl)chroman-6-yloxy]-5-nitropyridine**

15

a) 6-Hydroxy-2-(3-methoxyphenyl)chroman-4-one

6-Hydroxy-2-(3-methoxyphenyl)chroman-4-one was prepared as described for
6-hydroxy-2-(3-fluorophenyl)chroman-4-one in Example 9(a). The product was
20 recrystallised from ethanol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.42 (s, 1H), 7.33 (t,
1H, J 15.8, 8.3 Hz), 7.12 (d, 1H, J 3.0 Hz), 7.10 (s, 1H), 7.09 (d, 1H, J 8.3 Hz), 7.04
(dd, 1H, J 8.8, 3.0 Hz), 6.96 (d, 1H, 8.8 Hz), 6.93 (dd, 1H, J 8.0, 2.5 Hz), 5.52 (dd,
1H, J 12.9, 2.9 Hz), 3.77 (s, 3H), 3.17 (dd, 1H, J 16.9, 12.9 Hz), 2.77 (dd, 1H, J 16.9,
2.9 Hz).

25

b) 2-(3-methoxyphenyl)chroman-4,6-diol

2-(3-Methoxyphenyl)chroman-4,6-diol was prepared as described for 2-phenylchroman-4,6-diol in Example 8(a) starting from 6-hydroxy-2-(3-methoxyphenyl)-
30 chroman-4-one. ¹H NMR (400 MHz, d₆-DMSO) δ: 8.82 (s, 1H), 7.31 (t, 1H, J 15.7,
7.9 Hz), 6.99-7.02 (m, 2H), 6.88-6.90 (m, 2H), 6.59 (d, 1H, J 8.7 Hz), 6.54 (dd, 1H, J
8.7, 2.8 Hz), 5.40 (d, 1H, J 7.0 Hz), 5.08 (d, 1H, J 11.5 Hz), 4.83-4.89 (m, 1H), 3.77
(s, 3H), 2.23-2.28 (m, 1H), 1.83-1.92 (m, 1H).

35

c) 2-(3-Methoxyphenyl)chroman-6-ol

2-(3-Methoxyphenyl)chroman-6-ol was prepared as described for 2-(3-fluorophenyl)chroman-6-ol in Example 9(c) starting from 2-(3-methoxyphenyl)chroman-4,6-diol. ¹H NMR (300 MHz, d₆-DMSO) δ: 8.75 (s, 1H), 7.28 (t, 1H, J 15.7, 7.9 Hz), 6.96-6.99 (m, 2H), 6.87 (dd, 1H, J 7.9, 2.5 Hz), 6.63 (d, 1H, J 8.3 Hz), 6.52 (d, 1H, J 2.9 Hz), 6.48 (s, 1H), 4.95 (dd, 1H, J 9.8, 2.2 Hz), 3.75 (s, 3H), 2.82-2.89 (m, 1H), 2.57-2.66 (m, 1H), 2.06-2.13 (m, 1H), 1.89-1.97 (m, 1H).

d) 2-[2-(3-Methoxyphenyl)chroman-6-yloxy]-5-nitropyridine

2-[2-(3-Methoxyphenyl)chroman-6-yloxy]-5-nitropyridine was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(3-methoxyphenyl)chroman-6-ol. ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.60 (dd, 1H, J 9.1, 2.9 Hz), 7.32 (t, 1H, J 15.7, 7.9 Hz), 7.20 (d, 1H, J 9.1 Hz), 7.03 (d, 1H, J 8.4 Hz), 7.01 (s, 1H), 7.00 (d, 1H, J 2.8 Hz), 6.96 (dd, 1H, J 8.7, 2.8 Hz), 6.92 (d, 1H, J 8.7 Hz), 6.90 (dd, 1H, J 8.4, 2.6 Hz), 5.12 (dd, 1H, J 10.0, 2.3 Hz), 3.77 (s, 3H), 2.93-2.97 (m, 1H), 2.71-2.77 (m, 1H), 2.15-2.20 (m, 1H), 1.99-2.05 (m, 1H).

Example 100:

6-[2-(3-Methoxyphenyl)chroman-6-yloxy]pyridin-3-ylamine

6-[2-(3-Methoxyphenyl)chroman-6-yloxy]pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using 300 mg 2-[2-(3-methoxyphenyl)chroman-6-yloxy]-5-nitropyridine (Example 99(d)) and 1.0 g of Zn. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.51 (d, 1H, J 3.0 Hz), 7.31 (t, 1H, J 15.8, 7.9 Hz), 7.04 (dd, 1H, J 8.7, 3.0 Hz), 6.99-7.02 (m, 1H), 6.99 (d, 1H, J 2.6 Hz), 6.90 (dd, 1H, J 8.9, 2.6 Hz), 6.79-6.81 (m, 1H), 6.72-6.74 (m, 2H), 6.69 (d, 1H, J 8.9 Hz), 5.06 (dd, 1H, J 9.9, 2.2 Hz), 4.50 (s, 2H), 3.77 (s, 3H), 2.88-2.95 (m, 1H), 2.66-2.71 (m, 1H), 2.12-2.17 (m, 1H), 1.94-2.00 (m, 1H).

Example 101:

N-{6-[2-(3-Methoxyphenyl)chroman-6-yloxy]pyridin-3-yl}acetamide

N-{6-[2-(3-Methoxyphenyl)chroman-6-yloxy]pyridin-3-yl}acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 6-[2-(3-Methoxyphenyl)chroman-6-yloxy]pyridin-3-

ylamine (Example 100). ¹H NMR (400 MHz, d₆-DMSO) δ: 10.04 (s, 1H), 8.27 (d, 1H, J 2.7 Hz), 8.01 (dd, 1H, J 8.9, 2.7 Hz), 7.32 (t, 1H, J 15.7, 7.8 Hz), 7.02 (d, 1H, J 8.8 Hz), 7.00 (d, 1H, J 2.5 Hz), 6.92 (d, 1H, J 8.9 Hz), 6.90 (dd, 1H, J 8.2, 2.5 Hz), 6.84-6.86 (m, 3H), 5.09 (dd, 1H, J 9.9, 2.1 Hz), 3.77 (s, 3H), 2.91-2.95 (m, 1H),
5 2.68-2.74 (m, 1H), 2.14-2.18 (m, 1H), 2.04 (s, 3H), 1.97-2.02 (m, 1H).

Example 102:**2-(3-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol**

10 2-(3-Methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol was prepared as described for 5-nitro-2-(2-phenylchroman-6-yloxy)pyridine in Example 1(b) starting from 2-(3-methoxyphenyl)chroman-4,6-diol (Example 99(b)). ¹H NMR (400 MHz, d₆-DMSO) δ: 9.04 (d, 1H, J 2.9 Hz), 8.61 (dd, 1H, 9.1, 2.9 Hz), 7.34 (t, 1H, J 15.7, 7.8 Hz), 7.25 (d, 1H, J 2.4 Hz), 7.22 (d, 1H, 9.1 Hz), 7.00-7.06 (m, 3H), 6.92
15 (dd, 1H, J 8.8, 2.4 Hz), 6.88 (d, 1H, J 8.8 Hz), 5.64 (d, 1H, J 6.4 Hz), 5.27 (d, 1H, J 10.7 Hz), 4.95-5.00 (m, 1H), 3.79 (s, 3H), 2.30-2.35 (m, 1H), 1.92-2.01 (m, 1H).

Example 103:**6-(5-Aminopyridin-2-yloxy)-2-(3-methoxyphenyl)chroman-4-ol**

20 6-(5-Aminopyridin-2-yloxy)-2-(3-methoxyphenyl)chroman-4-ol was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 using 138 mg 2-(3-methoxyphenyl)-6-(5-nitropyridin-2-yloxy)chroman-4-ol (Example 102) and 457 mg of Zn. ¹H NMR (400 MHz, d₆-DMSO) δ: 7.52 (d, 1H, J 3.0 Hz), 7.32 (t, 1H, J 15.7, 7.8 Hz), 7.06 (dd, 1H, 8.7, 3.0 Hz), 7.01-7.04 (m, 3H), 6.91 (dd, 1H, J 8.6, 2.4 Hz), 6.78-6.80 (m, 1H), 6.76 (d, 1H, J 8.7 Hz), 6.71 (d, 1H, J 8.6 Hz), 5.54
25 (d, 1H, J 6.5 Hz), 5.19 (d, 1H, J 10.6 Hz), 5.01 (s, 2H), 4.88-4.94 (m, 1H), 3.78 (s, 3H), 2.27-2.32 (m, 1H), 1.87-1.96 (m, 1H).

Example 104:**6-(2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-ylamine hydrochloride**

35 6-(2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-ylamine was prepared as described for 5-amino-2-(2-phenylchroman-6-yloxy)pyridine in Example 26 starting from 5-nitro-2-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridine

(Example 84). 6-(2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-ylamine was isolated as its dihydrochloride salt. ¹H NMR (400 MHz, CDCl₃) δ: 8.20 (d, 1H, J 2.1 Hz), 7.87 (dd, 1H, J 8.9, 2.1 Hz), 7.41-7.44 (m, 4H), 7.37-7.40 (m, 1H), 6.98 (d, 1H, J 8.9 Hz), 6.97 (d, 1H, J 8.8 Hz), 6.93 (d, 1H, J 2.7 Hz), 6.80 (dd, 1H, J 8.8, 2.7 Hz), 5.20 (dd, 1H, J 9.6, 1.9 Hz), 3.30 (dd, 1H, 13.2, 9.6 Hz), 3.12 (dd, 1H, 13.2, 1.9 Hz).

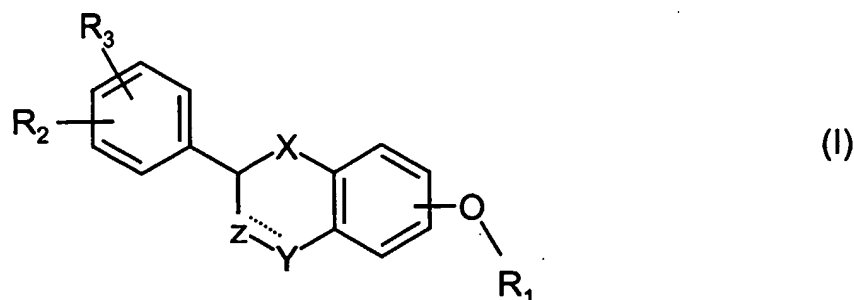
Example 105:

N-[6-(2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-yl]acet-
10 amide

N-[6-(2-Phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-yl]acetamide was prepared as described for 5-N'-acetylamino-2-(2-phenylchroman-6-yloxy)pyridine in Example 27 starting from 6-(2-phenyl-2,3-dihydrobenzo[1,4]oxathiin-6-yloxy)pyridin-3-ylamine. ¹H NMR (400 MHz, CDCl₃) δ: 8.07-8.11 (m, 2H), 7.40-7.42 (m, 4H), 7.36-7.39 (m, 1H), 7.15 (br s, 1H), 6.94 (d, 1H, J 8.8 Hz), 6.90 (d, 1H, J 2.8 Hz), 6.88 (d, 1H, J 9.1 Hz), 6.78 (dd, 1H, J 8.8, 2.8 Hz), 5.17 (dd, 1H, J 9.6, 1.9 Hz), 3.28 (dd, 1H, 13.2, 9.6 Hz), 3.08 (dd, 1H, 13.2, 1.9 Hz), 2.19 (s, 3H).

Claims

1. Compounds of formula (I):

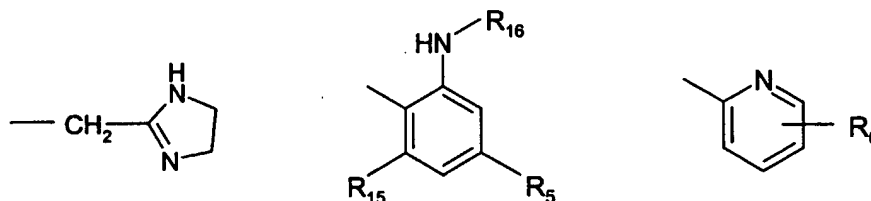


5

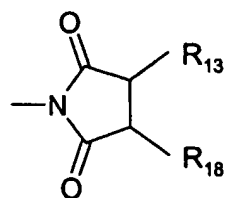
wherein

X is -O-, -CH₂- or -C(O)-;Z is -CHR₉- or valence bond;Y is -CH₂-, -C(O)-, CH(OR₁₀)-, -CH(NR₁₁R₁₂)-, -O-, -S-, -S(O)- or -S(O₂)-
provided that in case Z is a valence bond, Y is not C(O);

10

the dashed line represents an optional double bond in which case Z is -CR₉-
and Y is -CH-, C(OR₁₀)- or -C(NR₁₁R₁₂)-;R₁ is -(CH₂)_nNR₄R₇ or one of the following groups:

n is 1 - 4,

15 R₂ and R₃ are independently H, lower alkyl, lower alkoxy, -NO₂, halogen,
-CF₃, -OH, -NHR₈ or -COOH,R₄ and R₇ are independently H, lower alkyl or lower hydroxyalkyl,R₅ is H, lower alkoxy, -CF₃, -NH₂ or -CN,R₆ is -NO₂, -NR₁₄R₁₉, -CF₃ orR₈ and R₁₆ are independently H or acyl,

20

R₉ is H or lower alkyl,

R_{10} is H, alkylsulfonyl or acyl;

R_{11} and R_{12} are independently H, lower alkyl or acyl,

R_{13} and R_{18} are independently H or $-OR_{20}$,

R_{14} and R_{19} are independently H, acyl, alkylsulfonyl, $C(S)NHR_{17}$ or

5 $C(O)NHR_{17}$,

R_{15} is H or NH_2 ,

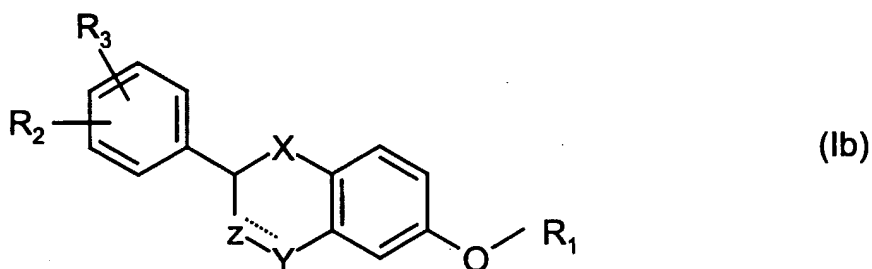
R_{17} is H or lower alkyl,

R_{20} is H or acyl,

and pharmaceutically acceptable salts and esters thereof.

10

2. A compound according to claim 1 having the structure



wherein R_2 , R_3 , X , Y and R_1 is as defined above.

15 3. A compound according to claim 1 or 2, wherein X is O, and Z and Y is $-CH_2-$.

4. A compound according to any of claims 1 or 2, wherein X is O, Z is $-CH_2$ and Y is $CHOH$.

20 5. A compound according to any of claims 1 to 4, wherein R_1 is one of the following groups:



6. A compound according to any of claims 1 to 5, wherein R_6 is NO_2 or $-NR_{14}R_{19}$.

25

7. A compound according to claim 6, wherein R_{14} and R_{19} are independently H, acyl or alkylsulfonyl.

8. A compound according to any of claims 1 to 7, wherein R_{15} and R_{16} is H.

5

9. A compound according to any of claims 1 to 8, wherein R_5 is H or lower alkoxy.

10. A compound according to any of claims 1 to 9, wherein R_2 and R_3 are independently H or halogen.

10

11. A compound according to claim 10, wherein halogen is fluorine.

12. A compound according to any of claims 1 to 11, wherein $n = 2$.

15

13. A compound according to any of claims 1 to 12, wherein R_4 and R_7 are methyl.

14. A pharmaceutical composition comprising a compound of claim 1 together with a pharmaceutically acceptable carrier.

20

15. A method for inhibiting Na^+/Ca^{2+} exchange mechanism in a cell, comprising administering to a subject in need thereof a therapeutically effective amount of a compound of claim 1.

25

16. A method for treating arrhythmias, comprising administering to a subject in need thereof a therapeutically effective amount of a compound of claim 1.

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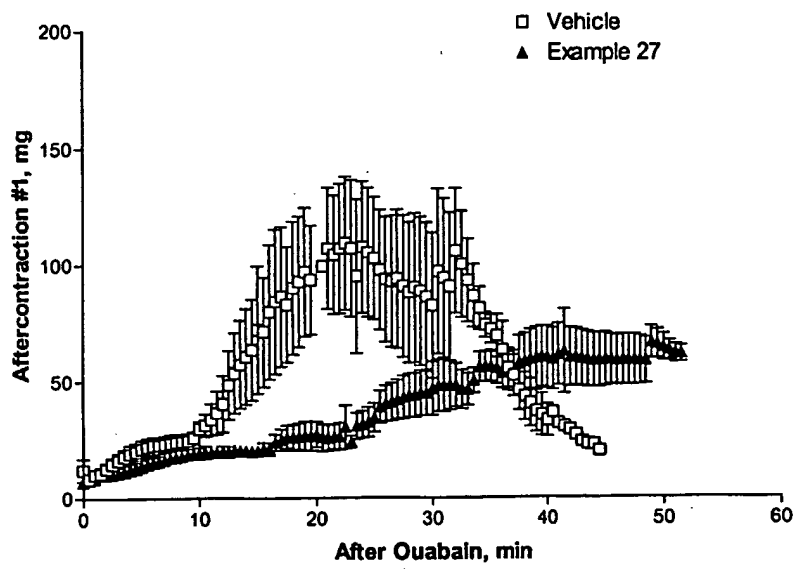


Fig. 1

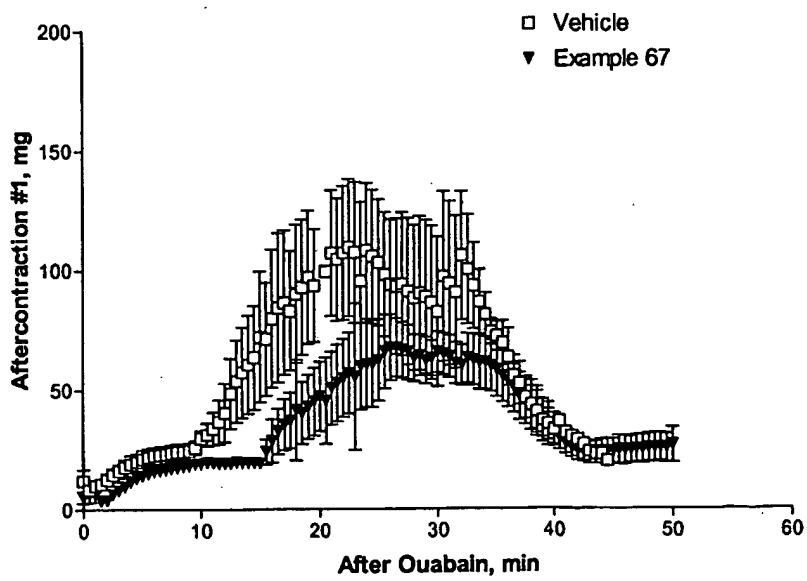


Fig. 2

INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 02/00621

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 311 /04 , 311 /32 , 319 /16 , 327 /06, 405 /12 , 411/12, C07C 217/14 , 217/54, 217/90,
A61K 31/135, 31/353, 31/357, 31/39, 31/4178, 31/4433, A61P 9/06, 9/10

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K, A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-INTERNAL, CHEM.ABS.DATA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 1154119 A (MERCK AKTIENGESSELLSCHAFT), 4 June 1969 (04.06.69) --	1,2,4,10,12, 13,14
X	GB 1068751 A (UPJOHN COMPANY), 10 May 1967 (10.05.67), see part. example 10 and claim 6 --	1-2,10,12,14
A	PATENT ABSTRACTS OF JAPAN vol.200, no. 002 29 February 2000 (2000-02-29) & JP 11 302235 A (TAISHO PHARMACEUT CO LTD), 02 November 1999 (1999-11-02) abstract --	1-16

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

23 October 2002

Date of mailing of the international search report

06. 11. 2002

Name and mailing address of the International Searching Authority



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INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 02/00621

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol.199, no. 905 31 May 1999 (1999-05-31) & JP 11 049752 A (TAISHO PHARMACEUT CO LTD), 23 February 1999 (1999-02-23) abstract --	1-16
A	EP 1031556 A1 (TAISHO PHARMACEUTICAL CO LTD), 30 August 2000 (30.08.00) --	1-16
A	US 5703118 A (DURAND ET AL), 30 December 1997 (30.12.97) --	1-16
A	WO 0064445 A1 (BRISTOL-MYERS SQUIBB COMPANY), 2 November 2000 (02.11.00) -- -----	1-16

INTERNATIONAL SEARCH REPORT

International application No.
PCT/FI02/00621**Box I** Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: **15, 16**
because they relate to subject matter not required to be searched by this Authority, namely:
see next sheet
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Claims 15,16 relate to methods of treatment of the human or animal body by surgery or by therapy (Rule. 39.1.(iv)). Nevertheless, a search has been executed for these claims. The search has been based on the vector used and on the alleged effects of the vector.

INTERNATIONAL SEARCH REPORT
Information on patent family members

30/09/02

International application No.

PCT/FI 02/00621

Patent document cited in search report			Publication date	Patent family member(s)		Publication date
GB	1154119	A	04/06/69	NONE		
GB	1068751	A	10/05/67	BE	650081 A	00/00/00
				CH	456579 A	31/07/68
				DE	1493323 A,B	17/07/69
				FR	1568704 A	30/05/69
				NL	6407603 A	04/01/65
				US	3862232 A	21/01/75
				US	3919327 A	11/11/75
EP	1031556	A1	30/08/00	AU	735185 B	05/07/01
				AU	9463898 A	10/05/99
				CA	2305734 A	29/04/99
				US	6162832 A	19/12/00
				CN	1276783 T	13/12/00
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US	5703118	A	30/12/97	AU	5204396 A	14/11/96
				CA	2175747 A	06/11/96
				CN	1136920 A	04/12/96
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				FR	2733685 A,B	08/11/96
				JP	8301754 A	19/11/96
				NO	961797 A	06/11/96
				ZA	9603520 A	13/11/96
WO	0064445	A1	02/11/00	AU	4360200 A	10/11/00
				EP	1180029 A	20/02/02